

General

Common Present on Admission Diagnosis

| | | |
|--------------------------|--|---------|
| <input type="checkbox"/> | Acidosis | Post-op |
| <input type="checkbox"/> | Acute Post-Hemorrhagic Anemia | Post-op |
| <input type="checkbox"/> | Acute Renal Failure | Post-op |
| <input type="checkbox"/> | Acute Respiratory Failure | Post-op |
| <input type="checkbox"/> | Acute Thromboembolism of Deep Veins of Lower Extremities | Post-op |
| <input type="checkbox"/> | Anemia | Post-op |
| <input type="checkbox"/> | Bacteremia | Post-op |
| <input type="checkbox"/> | Bipolar disorder, unspecified | Post-op |
| <input type="checkbox"/> | Cardiac Arrest | Post-op |
| <input type="checkbox"/> | Cardiac Dysrhythmia | Post-op |
| <input type="checkbox"/> | Cardiogenic Shock | Post-op |
| <input type="checkbox"/> | Decubitus Ulcer | Post-op |
| <input type="checkbox"/> | Dementia in Conditions Classified Elsewhere | Post-op |
| <input type="checkbox"/> | Disorder of Liver | Post-op |
| <input type="checkbox"/> | Electrolyte and Fluid Disorder | Post-op |
| <input type="checkbox"/> | Intestinal Infection due to Clostridium Difficile | Post-op |
| <input type="checkbox"/> | Methicillin Resistant Staphylococcus Aureus Infection | Post-op |
| <input type="checkbox"/> | Obstructive Chronic Bronchitis with Exacerbation | Post-op |
| <input type="checkbox"/> | Other Alteration of Consciousness | Post-op |
| <input type="checkbox"/> | Other and Unspecified Coagulation Defects | Post-op |
| <input type="checkbox"/> | Other Pulmonary Embolism and Infarction | Post-op |
| <input type="checkbox"/> | Phlebitis and Thrombophlebitis | Post-op |
| <input type="checkbox"/> | Protein-calorie Malnutrition | Post-op |
| <input type="checkbox"/> | Psychosis, unspecified psychosis type | Post-op |
| <input type="checkbox"/> | Schizophrenia Disorder | Post-op |
| <input type="checkbox"/> | Sepsis | Post-op |
| <input type="checkbox"/> | Septic Shock | Post-op |
| <input type="checkbox"/> | Septicemia | Post-op |
| <input type="checkbox"/> | Type II or Unspecified Type Diabetes Mellitus with Mention of Complication, Not Stated as Uncontrolled | Post-op |
| <input type="checkbox"/> | Urinary Tract Infection, Site Not Specified | Post-op |

Elective Outpatient, Observation, or Admission (Single Response)

| | | |
|--------------------------|---|--|
| <input type="checkbox"/> | Elective outpatient procedure: Discharge following routine recovery | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> | Outpatient observation services under general supervision | Diagnosis: Admitting Physician: Patient Condition: Bed request comments: PACU & Post-op |
| <input type="checkbox"/> | Outpatient in a bed - extended recovery | Diagnosis: Admitting Physician: Bed request comments: PACU & Post-op |
| <input type="checkbox"/> | Admit to Inpatient | Diagnosis: Admitting Physician: Level of Care: Patient Condition: Bed request comments: Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op |

Admission or Observation (Single Response)

Patient has active outpatient status order on file

- | | |
|--|--|
| <input type="checkbox"/> Admit to Inpatient | Diagnosis: Admitting Physician: Level of Care: Patient Condition: Bed request comments: Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op |
| <input type="checkbox"/> Outpatient observation services under general supervision | Diagnosis: Admitting Physician: Patient Condition: Bed request comments: PACU & Post-op |
| <input type="checkbox"/> Outpatient in a bed - extended recovery | Diagnosis: Admitting Physician: Bed request comments: PACU & Post-op |
| <input type="checkbox"/> Transfer patient | Level of Care: Bed request comments: Scheduling/ADT |
| <input type="checkbox"/> Return to previous bed | Routine, Until discontinued, Starting S, Scheduling/ADT |

Admission (Single Response)

Patient has active status order on file

- | | |
|---|--|
| <input type="checkbox"/> Admit to inpatient | Diagnosis: Admitting Physician: Level of Care: Patient Condition: Bed request comments: Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op |
| <input type="checkbox"/> Transfer patient | Level of Care: Bed request comments: Scheduling/ADT |
| <input type="checkbox"/> Return to previous bed | Routine, Until discontinued, Starting S, Scheduling/ADT |

Transfer (Single Response)

Patient has active inpatient status order on file

- | | |
|---|---|
| <input type="checkbox"/> Transfer patient | Level of Care: Bed request comments: Scheduling/ADT |
| <input type="checkbox"/> Return to previous bed | Routine, Until discontinued, Starting S, Scheduling/ADT |

Code Status

- | | |
|---|--|
| <input type="checkbox"/> Full Code | Code Status decision reached by: Post-op |
| <input type="checkbox"/> DNR (Do Not Resuscitate) | |
| <input type="checkbox"/> DNR (Do Not Resuscitate) | Does patient have decision-making capacity? Post-op |

| | |
|---|--|
| <input type="checkbox"/> Consult to Palliative Care Service | Priority: Reason for Consult? Order? Name of referring provider: Enter call back number: |
| <input type="checkbox"/> Consult to Social Work | Reason for Consult: Post-op |
| <input type="checkbox"/> Modified Code | Does patient have decision-making capacity? Modified Code restrictions: Post-op |
| <input type="checkbox"/> Treatment Restrictions | Treatment Restriction decision reached by: Specify Treatment Restrictions: Post-op |

Isolation

| | |
|--|---------|
| <input type="checkbox"/> Airborne isolation status | Details |
| <input type="checkbox"/> Contact isolation status | Details |
| <input type="checkbox"/> Droplet isolation status | Details |
| <input type="checkbox"/> Enteric isolation status | Details |

Precautions

| | |
|---|--|
| <input type="checkbox"/> Aspiration precautions | Post-op |
| <input type="checkbox"/> Fall precautions | Increased observation level needed: Post-op |
| <input type="checkbox"/> Latex precautions | Post-op |
| <input type="checkbox"/> Seizure precautions | Increased observation level needed: Post-op |

Nursing

Discharge Instructions

| | |
|---|-----------------------|
| <input checked="" type="checkbox"/> Follow-up with provider | Provider: On date: |
| <input type="checkbox"/> Follow-up with primary care provider | Routine |
| <input type="checkbox"/> Ambulate patient | Routine |
| <input type="checkbox"/> Patient may shower daily | Routine |
| <input type="checkbox"/> No heavy lifting greater than 15 lbs | Routine |
| <input type="checkbox"/> No driving for 2 weeks | Routine |
| <input type="checkbox"/> If patient discharged with foley catheter, provide teaching regarding care of leg bag and bedside bag. | Routine |

Vital Signs

| | |
|---|--|
| <input checked="" type="checkbox"/> Vital signs - T/P/R/BP - per recovery routine | Routine, Per unit protocol, PACU & Post-op |
|---|--|

Nursing Interventions

| | |
|--|---|
| <input type="checkbox"/> Ambulate | Routine, As needed Specify: with assistance PACU |
| <input type="checkbox"/> Foley catheter care | Routine, Until discontinued, Starting S Orders: Maintain PACU |

| | |
|---|---|
| <input type="checkbox"/> Foley catheter - discontinue | <p>Routine, Once If the urine is clear and a post void residual (PVR) checked.</p> <p>If PVR less than 150 mL may discharge and if PVR greater than 200 mL, notify MD. PACU Cont. To floor/ICU, Patient due to void within 6 hours of catheter removal.</p> <p>- If patient voids more than 200 mL -- continue to monitor I&O - If patient voids less than 200 mL within 6 hours after catheter removal, is uncomfortable or incontinent - perform bladder scan to assess residual volume.</p> <p>Follow nurse driven catheter removal algorithm if scanned volume greater than 200 mL: straight cath and record residual. Re-evaluate in 6 hours or sooner if symptomatic. May repeat x1. Scanned volume greater than 200 mL after straight cathed twice, contact physician.</p> <p>Catheter use: does not meet CDC criteria for catheter use., PACU</p> |
| <input type="checkbox"/> Patient education (specify) | <p>Routine, Once Patient/Family: Education for: If discharging with foley catheter, provide teaching regarding care of leg bag and bedside bag., PACU</p> |
| <input type="checkbox"/> If Mitomycin C in the bladder: | <p>Routine, Until discontinued, Starting S Leave mitomycin C indwelling for at least 30 minutes - up to 60 minutes. Rotate patient from side to side every 20 minutes; then drain bladder, avoiding contact with mitomycin C and remove catheter., PACU</p> |

IV Fluids

IV Fluids (Single Response)

| | |
|--|---|
| <input type="checkbox"/> sodium chloride 0.9 % infusion | 100 mL/hr, intravenous, continuous, Post-op |
| <input type="checkbox"/> lactated Ringer's infusion | 100 mL/hr, intravenous, continuous, Post-op |
| <input type="checkbox"/> dextrose 5%-0.9% sodium chloride infusion | 100 mL/hr, intravenous, continuous, Post-op |
| <input type="checkbox"/> dextrose 5 % and sodium chloride 0.9 % with potassium chloride 20 mEq/L infusion | 100 mL/hr, intravenous, at 100 mL/hr, continuous, Post-op |
| <input type="checkbox"/> dextrose 5 % and lactated Ringer's infusion | 100 mL/hr, intravenous, continuous, Post-op |
| <input type="checkbox"/> dextrose 5%-0.45% sodium chloride infusion | 100 mL/hr, intravenous, continuous, Post-op |
| <input type="checkbox"/> dextrose 5 % and sodium chloride 0.45 % with potassium chloride 20 mEq/L infusion | 100 mL/hr, intravenous, continuous, Post-op |

Medications

Medications

| | |
|---|--|
| <input type="checkbox"/> oxybutynin (DITROPAN) tablet | 5 mg, oral, once PRN, bladder spasms, For 1 Doses, PACU & Post-op PACU continue to Floor/ICU |
| <input type="checkbox"/> belladonna alkaloids-opium (B&O SUPPRETTES) 16.2-30 mg suppository | 30 mg, rectal, once PRN, bladder spasms, For 1 Doses, PACU & Post-op Use if patient cannot take oral medication or if oral medication is ineffective. PACU continue to Floor/ICU |
| <input type="checkbox"/> phenazopyridine (PYRIDIDIUM) tablet | 200 mg, oral, once PRN, dysuria, For 1 Doses, PACU & Post-op PACU continue to ICU/Floor |
| <input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 7.5-325 mg per tablet | 1 tablet, oral, once PRN, moderate pain (score 4-6), For 1 Doses, PACU & Post-op |

Medications- HMSTJ

| | |
|---|--|
| <input type="checkbox"/> oxybutynin (DITROPAN) tablet | 5 mg, oral, once PRN, bladder spasms, For 1 Doses, PACU & Post-op PACU continue to Floor/ICU |
| <input type="checkbox"/> belladonna alkaloids-opium (B&O SUPPRETTES) 16.2-60 MG suppository | 60 mg, rectal, once PRN, bladder spasms, For 1 Doses, PACU & Post-op Use if patient cannot take oral medication or if oral medication is ineffective. PACU continue to Floor/ICU |
| <input type="checkbox"/> phenazopyridine (PYRIDIUM) tablet | 200 mg, oral, once PRN, dysuria, For 1 Doses, PACU & Post-op PACU continue to ICU/Floor |
| <input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 7.5-325 mg per tablet | 1 tablet, oral, once PRN, moderate pain (score 4-6), For 1 Doses, PACU & Post-op |

Antiemetics - HHM, HMSJ, HMW, HMSTC, HMTW Only

| | |
|---|--|
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral | "Or" Linked Panel |
| <input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet | 4 mg, oral, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection | 4 mg, intravenous, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is UNable to tolerate oral medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) IV or Oral or Rectal | "Or" Linked Panel |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) 12.5 mg IV | 12.5 mg, intravenous, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) tablet | 12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) suppository | 12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication. |

Antiemetics - HMSL, HMWB Only

| | |
|--|--|
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral | "Or" Linked Panel |
| <input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet | 4 mg, oral, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection | 4 mg, intravenous, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is UNable to tolerate oral medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) IV or Oral or Rectal | "Or" Linked Panel |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) 12.5 mg in sodium chloride 0.9 % 0.9 % 20 mL for Alaris pump syringe option | 12.5 mg, intravenous, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) tablet | 12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) suppository | 12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication. |

Antiemetics - HMSTJ Only

| | |
|--|--------------------------|
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral | "Or" Linked Panel |
|--|--------------------------|

| | |
|--|--|
| <input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet | 4 mg, oral, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection | 4 mg, intravenous, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is UNable to tolerate oral medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) IVPB or Oral or Rectal | "Or" Linked Panel |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) 25 mg in sodium chloride 0.9 % 50 mL IVPB | 12.5 mg, intravenous, for 30 Minutes, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) tablet | 12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is able to tolerate oral medication. |
| <input checked="" type="checkbox"/> promethazine (PHENERGAN) suppository | 12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication. |

Bowel Care (Single Response)

| | |
|---|---|
| <input type="checkbox"/> sennosides-docusate sodium (SENOKOT-S) 8.6-50 mg per tablet | 2 tablet, oral, nightly PRN, constipation, Post-op |
| <input type="checkbox"/> simethicone (MYLICON) chewable tablet | 160 mg, oral, 4 times daily PRN, flatulence, Post-op |
| <input type="checkbox"/> docusate sodium (COLACE) capsule | 100 mg, oral, 2 times daily PRN, constipation, Post-op |
| <input type="checkbox"/> magnesium hydroxide suspension - NOT RECOMMENDED FOR CHRONIC KIDNEY DISEASE STAGE 3 OR GREATER | 30 mL, oral, every 12 hours PRN, constipation, Post-op Do not give if patient is on hemodialysis or is in chronic renal failure. |
| <input type="checkbox"/> bisacodyl (DULCOLAX) EC tablet | 10 mg, oral, daily PRN, constipation, Post-op |
| <input type="checkbox"/> bisacodyl (DULCOLAX) suppository | 10 mg, rectal, daily PRN, constipation, Post-op |

Itching: For Patients GREATER than 77 years old (Single Response)

| | |
|---|---|
| <input type="checkbox"/> cetirizine (ZyrTEC) tablet | 5 mg, oral, daily PRN, itching, Post-op |
|---|---|

Itching: For Patients between 70-76 years old (Single Response)

| | |
|---|---|
| <input type="checkbox"/> cetirizine (ZyrTEC) tablet | 5 mg, oral, daily PRN, itching, Post-op |
|---|---|

Itching: For Patients LESS than 70 years old (Single Response)

| | |
|---|--|
| <input type="checkbox"/> diphenhydrAMINE (BENADRYL) tablet | 25 mg, oral, every 6 hours PRN, itching, Post-op |
| <input type="checkbox"/> hydrOXYzine (ATARAX) tablet | 10 mg, oral, every 6 hours PRN, itching, Post-op |
| <input type="checkbox"/> cetirizine (ZyrTEC) tablet | 5 mg, oral, daily PRN, itching, Post-op |
| <input type="checkbox"/> fexofenadine (ALLEGRA) tablet - For eGFR LESS than 80 mL/min, reduce frequency to once daily as needed | 60 mg, oral, 2 times daily PRN, itching, Post-op |

Insomnia: For Patients GREATER than 70 years old (Single Response)

| | |
|---|---|
| <input type="checkbox"/> ramelteon (ROZEREM) tablet | 8 mg, oral, nightly PRN, sleep, Post-op |
|---|---|

Insomnia: For Patients LESS than 70 years old (Single Response)

| | |
|---|---|
| <input type="checkbox"/> zolpidem (AMBIEN) tablet | 5 mg, oral, nightly PRN, sleep, Post-op |
| <input type="checkbox"/> ramelteon (ROZEREM) tablet | 8 mg, oral, nightly PRN, sleep, Post-op |

VTE

DVT Risk and Prophylaxis Tool (Single Response)

Low Risk Definition Moderate Risk Definition

Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. High Risk Definition

Both pharmacologic AND mechanical prophylaxis must be addressed.

Age less than 60 years and NO other VTE risk factors One or more of the following medical conditions: One or more of the following medical conditions:

Patient already adequately anticoagulated CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders)

Age 60 and above Severe fracture of hip, pelvis or leg

Central line Acute spinal cord injury with paresis

History of DVT or family history of VTE Multiple major traumas

Anticipated length of stay GREATER than 48 hours Abdominal or pelvic surgery for CANCER

Less than fully and independently ambulatory Acute ischemic stroke

Estrogen therapy History of PE

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Low Risk of DVT

Low Risk (Single Response)

Low risk of VTE

Routine, Once

Low risk: Due to low risk, no VTE prophylaxis is needed.

Will encourage early ambulation

PACU & Post-op

Moderate Risk of DVT - Surgical

Address pharmacologic prophylaxis by selecting one of the following. Mechanical prophylaxis is optional unless pharmacologic prophylaxis is contraindicated.

Moderate Risk

Moderate risk of VTE

Routine, Once, PACU & Post-op

Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response)

Patient is currently receiving therapeutic anticoagulation

Routine, Once

No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication.

Therapy for the following:

PACU & Post-op

Contraindications exist for pharmacologic prophylaxis

Routine, Once

No pharmacologic VTE prophylaxis due to the following contraindication(s):

PACU & Post-op

enoxaparin (LOVENOX) injection (Single Response)

enoxaparin (LOVENOX) syringe

40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1

enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min

30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1

For Patients with CrCL LESS than 30 mL/min

enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min

30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1

For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min

enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min

40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1

For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min

| | |
|--|--|
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk of DVT - Non-Surgical | |
| Address pharmacologic prophylaxis by selecting one of the following. Mechanical prophylaxis is optional unless pharmacologic prophylaxis is contraindicated. | |
| <input type="checkbox"/> Moderate Risk | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |

| | |
|--|--|
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT), do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk of DVT - Surgical Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis. | |
| <input type="checkbox"/> High Risk | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCl LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCl LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |

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| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk of DVT - Non-Surgical | |
| Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis. | |
| <input type="checkbox"/> High Risk | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCl LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCl LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |

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| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis. | |
| <input type="checkbox"/> High Risk | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> apixaban (ELIQUIS) tablet | 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: |
| <input type="checkbox"/> aspirin chewable tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| <input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - hip arthroplasty | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - knee arthroplasty | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min - knee/hip arthroplasty | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min. |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. |

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| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (time critical), Starting S+1, PACU & Post-op To be Given on Post Op Day 1. Indications: |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |

DVT Risk and Prophylaxis Tool (Single Response)

Low Risk Definition Moderate Risk Definition

Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. High Risk Definition

Both pharmacologic AND mechanical prophylaxis must be addressed.

Age less than 60 years and NO other VTE risk factors One or more of the following medical conditions: One or more of the following medical conditions:

Patient already adequately anticoagulated CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders)

Age 60 and above Severe fracture of hip, pelvis or leg

Central line Acute spinal cord injury with paresis

History of DVT or family history of VTE Multiple major traumas

Anticipated length of stay GREATER than 48 hours Abdominal or pelvic surgery for CANCER

Less than fully and independently ambulatory Acute ischemic stroke

Estrogen therapy History of PE

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Low Risk of DVT

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| <input type="checkbox"/> Low Risk (Single Response) | |
| <input type="checkbox"/> Low risk of VTE | Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation PACU & Post-op |
| <input type="checkbox"/> Moderate Risk of DVT - Surgical | |
| Address pharmacologic prophylaxis by selecting one of the following. Mechanical prophylaxis is optional unless pharmacologic prophylaxis is contraindicated. | |
| <input type="checkbox"/> Moderate Risk | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |

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| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk of DVT - Non-Surgical | |
| Address pharmacologic prophylaxis by selecting one of the following. Mechanical prophylaxis is optional unless pharmacologic prophylaxis is contraindicated. | |
| <input type="checkbox"/> Moderate Risk | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT), do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |

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| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk of DVT - Surgical | |
| Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis. | |
| <input type="checkbox"/> High Risk | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |

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| <input type="checkbox"/> | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> | Place antiembolic stockings | Routine, Once, PACU & Post-op |
| () High Risk of DVT - Non-Surgical | | |
| Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis. | | |
| [] High Risk | | |
| <input type="checkbox"/> | High risk of VTE | Routine, Once, PACU & Post-op |
| [] High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | | |
| <input type="checkbox"/> | Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> | Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () enoxaparin (LOVENOX) injection (Single Response) | | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> | fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> | heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, PACU & Post-op |
| <input type="checkbox"/> | heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> | warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| <input type="checkbox"/> | Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | | |
| <input type="checkbox"/> | Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> | Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> | Place antiembolic stockings | Routine, Once, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) | | |

Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.

| | |
|--|--|
| <input type="checkbox"/> High Risk | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) | |
| <input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> apixaban (ELIQUIS) tablet | 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: |
| <input type="checkbox"/> aspirin chewable tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| <input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - hip arthroplasty | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - knee arthroplasty | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min - knee/hip arthroplasty | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min. |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| <input type="checkbox"/> fondaparinux (ARIXTRA) injection | 2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT): |
| <input type="checkbox"/> heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| <input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| <input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (time critical), Starting S+1, PACU & Post-op To be Given on Post Op Day 1. Indications: |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| <input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| <input type="checkbox"/> Mechanical Prophylaxis (Single Response) | |

| | |
|---|---|
| <input type="checkbox"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place sequential compression device and antiembolic stockings | "And" Linked Panel |
| <input type="checkbox"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings | Routine, Once, PACU & Post-op |

Labs

Labs

| | |
|--|------------|
| <input type="checkbox"/> Hemoglobin and hematocrit | Once, PACU |
|--|------------|

Cardiology

Imaging

Other Studies

Respiratory

Rehab

Consults

For Physician Consult orders use sidebar

Additional Orders