

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="radio"/> Elective outpatient procedure: Discharge following routine recovery | Routine, Continuous, PACU & Post-op |
| <input type="radio"/> Outpatient observation services under general supervision | Admitting Physician: Patient Condition: Bed request comments: PACU & Post-op |
| <input type="radio"/> Outpatient in a bed - extended recovery | Admitting Physician: Bed request comments: PACU & Post-op |
| <input type="radio"/> Admit to Inpatient | 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 | 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 | Admitting Physician: Patient Condition: Bed request comments: PACU & Post-op |
| <input type="checkbox"/> Outpatient in a bed - extended recovery | 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 | 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) (Selection Required) | |
| <input type="checkbox"/> DNR (Do Not Resuscitate) | Did the patient/surrogate require the use of an interpreter? Did the patient/surrogate require the use of an interpreter? 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 | Did the patient/surrogate require the use of an interpreter? Did the patient/surrogate require the use of an interpreter? Does patient have decision-making capacity? Modified Code restrictions: Post-op |
| <input type="checkbox"/> Treatment Restrictions | I understand that if the patient is NOT in a cardiopulmonary arrest, the selected treatments will NOT be provided. I understand that all other unselected medically indicated treatments will be provided. Treatment Restriction decision reached by: Specify Treatment Restrictions: Post-op |

Isolation

| | |
|---|-----------------------|
| <input type="checkbox"/> Airborne isolation status | |
| <input type="checkbox"/> Airborne isolation status | Details |
| <input type="checkbox"/> Mycobacterium tuberculosis by PCR - If you suspect Tuberculosis, please order this test for rapid diagnostics. | Once, Sputum, Post-op |
| <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 | PACU & Post-op |
| <input checked="" type="checkbox"/> Fall precautions | Increased observation level needed: PACU & Post-op |
| <input type="checkbox"/> Latex precautions | PACU & Post-op |
| <input type="checkbox"/> Seizure precautions | Increased observation level needed: PACU & Post-op |
| <input type="checkbox"/> Spinal precautions | PACU & Post-op |

Nursing

Vital Signs (Single Response)

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

Activity

| | |
|---|---|
| <input type="checkbox"/> Strict bed rest | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input type="checkbox"/> Up with assistance | Routine, Until discontinued, Starting S Specify: Up with assistance PACU & Post-op |
| <input type="checkbox"/> Activity as tolerated | Routine, Until discontinued, Starting S Specify: Activity as tolerated PACU & Post-op |
| <input type="checkbox"/> Ambulate with assistance | Routine, Every 8 hours Specify: with assistance, in hall Ambulate with assistance at least 3 times a day, ambulate in hallway by tomorrow AM., PACU & Post-op |
| <input type="checkbox"/> Up in chair for all meals | Routine, Until discontinued, Starting S Specify: Up in chair Additional modifier: for meals All meals, PACU & Post-op |
| <input type="checkbox"/> Elevate Head of bed 30 degrees | Routine, Until discontinued, Starting S Head of bed: 30 degrees PACU & Post-op |
| <input type="checkbox"/> Head of bed flat | Routine, Until discontinued, Starting S Head of bed: flat PACU & Post-op |

Nursing

| | |
|---|---|
| <input checked="" type="checkbox"/> Assess for neck swelling and airway compromise | Routine, Every 4 hours For 24 Hours Assess: for neck swelling and airway compromise PACU & Post-op |
| <input checked="" type="checkbox"/> Straight cath | Routine, Every 6 hours If unable to void after second attempt, insert Foley and call physician., PACU & Post-op |
| <input checked="" type="checkbox"/> Insert/Maintain Foley and Notify | |
| <input checked="" type="checkbox"/> Insert Foley catheter | Routine, Once Type: Size: Urinometer needed: If unable to void after second attempt at straight cath, insert Foley and call physician, PACU & Post-op |
| <input checked="" type="checkbox"/> Foley catheter care | Routine, Until discontinued, Starting S Orders: Maintain to gravity/bedside drain, PACU & Post-op |
| <input checked="" type="checkbox"/> Notify Physician if unable to void after second attempt at straight cath and Foley inserted | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input type="checkbox"/> Surgical/incision site care | Routine, Once Location: Site: Apply: Dressing Type: Open to air? PACU & Post-op |
| <input type="checkbox"/> Reinforce dressing | Routine, As needed Reinforce with: If saturated., PACU & Post-op |
| <input type="checkbox"/> Cervical collar - soft | Routine, Once Type of Collar to Apply: Soft cervical collar Special Instructions: obtain from central supply PACU & Post-op |
| <input type="checkbox"/> Cervical collar - Philadelphia | Routine, Once Type of Collar to Apply: Philadelphia Collar Special Instructions: Obtain from central supply. PACU & Post-op |
| <input type="checkbox"/> Cervical collar - Miami J | Routine, Once Type of Collar to Apply: Miami J Collar Special Instructions: Obtain from orthotic provider. PACU & Post-op |
| <input type="checkbox"/> Call Raborn Orthotics at 713-349-8117 for application of orthotic device | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input type="checkbox"/> Drain care | Routine, Until discontinued, Starting S Drain 1: Drain 2: Drain 3: Drain 4: All Drains: PACU & Post-op |
| <input type="checkbox"/> Place antiembolic stockings - Bilateral thigh high | Routine, Once May remove once patient ambulatory, PACU & Post-op |
| <input checked="" type="checkbox"/> No anticoagulants INcluding UNfractionated heparin | Routine, Until discontinued, Starting S Reason for "No" order: Postop Cervical Fusion PACU & Post-op |
| <input checked="" type="checkbox"/> No anti-platelet agents INcluding aspirin | Routine, Until discontinued, Starting S Reason for "No" order: Postop Cervical Fusion PACU & Post-op |

Notify (Selection Required)

| | |
|--|---|
| <input checked="" type="checkbox"/> Notify Physician of neck swelling or if airway compromised | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input checked="" type="checkbox"/> Notify Physician if acute change in neurological status | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input checked="" type="checkbox"/> Notify Physician bleeding at site | Routine, Until discontinued, Starting S, PACU & Post-op |
| <input checked="" type="checkbox"/> Notify Physician of No Bowel Movement for more than 72 hours | Routine, Until discontinued, Starting S, PACU & Post-op |

Diet

| | |
|--|--|
| <input checked="" type="checkbox"/> Diet - Clear liquids (advance as tolerated to Regular) | Diet effective now, Starting S Diet(s): Clear Liquids Advance Diet as Tolerated? Yes Target Diet: Regular Advance target diet criteria: Please assess bowel sounds between progressions. IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: PACU & Post-op |
| <input type="checkbox"/> Diet - Full liquids | Diet effective now, Starting S Diet(s): Full Liquids Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: PACU & Post-op |
| <input type="checkbox"/> Diet - Regular | Diet effective now, Starting S Diet(s): Regular Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: PACU & Post-op |
| <input type="checkbox"/> Diet - Heart healthy | Diet effective now, Starting S Diet(s): Heart Healthy Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: PACU & Post-op |
| <input type="checkbox"/> Diet - 2000 Kcal/255 gm Carb | Diet effective now, Starting S Diet(s): 2000 Kcal/225 gm Carbohydrate Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: PACU & Post-op |
| <input type="checkbox"/> Diet | Diet effective now, Starting S Diet(s): Other Options: Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid: Foods to Avoid: PACU & Post-op |

Education

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|---|---|
| <input type="checkbox"/> Patient education - Activity | Routine, Once Patient/Family: Education for: Activity PACU & Post-op |
|---|---|

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|---|--|
| <input checked="" type="checkbox"/> Patient education - Deep breathing and coughing exercises | Routine, Once Patient/Family: Education for: Other (specify) Specify: Deep breathing and coughing exercises PACU & Post-op |
| <input checked="" type="checkbox"/> Patient education - Incentive spirometry | Routine, Once Patient/Family: Education for: Incentive spirometry PACU & Post-op |
| <input checked="" type="checkbox"/> Patient education - Pain management | Routine, Once Patient/Family: Education for: Other (specify) Specify: Pain management PACU & Post-op |
| <input type="checkbox"/> Patient education - Smoking cessation | Routine, Once Patient/Family: Education for: Smoking cessation counseling PACU & Post-op |
| <input checked="" type="checkbox"/> Patient education - Wound care | Routine, Once Patient/Family: Education for: Other (specify) Specify: Wound care PACU & Post-op |

IV Fluids

IV Fluids (Single Response)

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

Medications

Steroids (Single Response)

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|--|--|
| <input type="checkbox"/> dexamethasone (DECADRON) IV | 4 mg, intravenous, every 6 hours scheduled, Post-op |
| <input type="checkbox"/> methylPREDNISolone sodium succinate (Solu-MEDROL) injection | 40 mg, intravenous, every 6 hours scheduled, Post-op |
| <input type="checkbox"/> methylPREDNISolone (MEDROL PAK) dose pack (start in AM) | |

THIS A PANEL. DO NOT EDIT.

| | |
|---|--|
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 8 mg, oral, before breakfast - one time, For 1 Doses, Post-op |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 4 mg, oral, after lunch - one time, S at 12:00 PM, For 1 Doses, Post-op |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 4 mg, oral, after dinner - one time, For 1 Doses, Post-op All day-1 doses may be given (up to 6 tablets) may be given at one time based on time of day. |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 8 mg, oral, nightly - one time, For 1 Doses, Post-op All day-1 doses may be given (up to 6 tablets) may be given at one time based on time of day. |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 4 mg, oral, 3 times daily around food, Starting S+1, For 3 Doses, Post-op |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 8 mg, oral, nightly - one time, Starting S+1, For 1 Doses, Post-op |
| <input type="checkbox"/> methylPREDNISolone (MEDROL) tablet | 4 mg, oral, 4 times daily tapering, Starting S+2, Post-op |

NSAIDS (Single Response)

These orders should be used for Cervical Arthroplasty. Use in spinal fusion patients is not recommended.

| | |
|--|---|
| <input type="checkbox"/> indomethacin (INDOCIN) capsule | 50 mg, oral, 3 times daily with meals, Starting S+1, Post-op Start POD #1. Use in spinal fusion patients not recommended. |
| <input type="checkbox"/> indomethacin SR (INDOCIN SR) CR capsule | 75 mg, oral, daily with breakfast, Starting S+1, Post-op Start POD #1. Use in spinal fusion patients not recommended. |

Medications

| | |
|---|--|
| <input type="checkbox"/> pantoprazole (PROTONIX) IV or ORAL | "Or" Linked Panel |
| <input type="checkbox"/> pantoprazole (PROTONIX) EC tablet | 40 mg, oral, daily at 0600, Post-op Indication(s) for Proton Pump Inhibitor (PPI) Therapy: |
| <input type="checkbox"/> pantoprazole (PROTONIX) 40 mg in sodium chloride 0.9 % 10 mL injection | 40 mg, intravenous, daily at 0600, Post-op Indication(s) for Proton Pump Inhibitor (PPI) Therapy: |

Medications - Bowel Management

| | |
|--|--------------------------------------|
| <input checked="" type="checkbox"/> polyethylene glycol (MIRALAX) packet | 17 g, oral, 2 times daily, Post-op |
| <input checked="" type="checkbox"/> Stool Softener Options (Single Response) | |
| <input checked="" type="checkbox"/> docusate sodium (COLACE) capsule | 100 mg, oral, 2 times daily, Post-op |
| <input type="checkbox"/> sennosides-docusate sodium (SENOKOT-S) 8.6-50 mg per tablet | 2 tablet, oral, nightly, Post-op |

Antibiotics (Single Response)

| | |
|--|---|
| <input type="checkbox"/> Antibiotics - Neurosurgery - patients with surgical site drains | |
| <input type="checkbox"/> Antibiotics: For Patients LESS than or EQUAL to 120 kg | |
| <input type="checkbox"/> cefazolin (ANCEF) IV - until drains removed | 1 g, intravenous, every 8 hours Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> cefepime (MAXIPIME) IV | 2 g, intravenous, every 12 hours Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> vancomycin (VANCOCIN) | 15 mg/kg, intravenous, once, For 1 Doses On call to cath lab. Transport antibiotics with patient to cath lab and administer prior to start of procedure Reason for Therapy: Surgical Prophylaxis |
| <input type="checkbox"/> Pharmacy consult to manage vancomycin | STAT, Until discontinued, Starting S Indication: Implanted Device Prophylaxis Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration Duration of Therapy (Days): |
| <input type="checkbox"/> Antibiotics: For Patients GREATER than 120 kg | |
| <input type="checkbox"/> cefazolin (ANCEF) IV - until drains removed | 1 g, intravenous, every 8 hours Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> cefepime (MAXIPIME) IV | 2 g, intravenous, every 12 hours Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> vancomycin (VANCOCIN) | 15 mg/kg, intravenous, once, For 1 Doses On call to cath lab. Transport antibiotics with patient to cath lab and administer prior to start of procedure Reason for Therapy: Surgical Prophylaxis |

| | | |
|---|---|---|
| <input type="checkbox"/> | Pharmacy consult to manage vancomycin | STAT, Until discontinued, Starting S Indication: Implanted Device Prophylaxis Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration Duration of Therapy (Days): |
| () Antibiotics - Neurosurgery - patients withOUT surgical site drains | | |
| <input type="checkbox"/> Antibiotics: For Patients LESS than or EQUAL to 120 kg | | |
| <input type="checkbox"/> | cefazolin (ANCEF) IV - until drains removed | 2 g, intravenous, once, For 1 Doses Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> | cefepime (MAXIPIME) IV | 2 g, intravenous, once, For 1 Doses Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required) | | |
| <input type="checkbox"/> | vancomycin (VANCOGIN) | 15 mg/kg, intravenous, once, For 1 Doses On call to cath lab. Transport antibiotics with patient to cath lab and administer prior to start of procedure Reason for Therapy: Surgical Prophylaxis |
| <input type="checkbox"/> | Pharmacy consult to manage vancomycin | STAT, Until discontinued, Starting S Indication: Implanted Device Prophylaxis Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration Duration of Therapy (Days): |
| <input type="checkbox"/> Antibiotics: For Patients GREATER than 120 kg | | |
| <input type="checkbox"/> | cefazolin (ANCEF) IV - until drains removed | 2 g, intravenous, once, For 1 Doses Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> | cefepime (MAXIPIME) IV | 2 g, intravenous, once, For 1 Doses Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration |
| <input type="checkbox"/> vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required) | | |
| <input type="checkbox"/> | vancomycin (VANCOGIN) | 15 mg/kg, intravenous, once, For 1 Doses On call to cath lab. Transport antibiotics with patient to cath lab and administer prior to start of procedure Reason for Therapy: Surgical Prophylaxis |
| <input type="checkbox"/> | Pharmacy consult to manage vancomycin | STAT, Until discontinued, Starting S Indication: Implanted Device Prophylaxis Reason for Therapy: Surgical Prophylaxis Surgical Prophylaxis: Please follow institutional and service line-specific guidelines for surgical prophylaxis for the stop date/duration Duration of Therapy (Days): |

Medications

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|--------------------------|---|--|
| <input type="checkbox"/> | benzocaine-menthol (CEPACOL MAX) lozenge 15-3.6 mg | 1 lozenge, buccal, PRN, sore throat, Post-op |
| <input type="checkbox"/> | phenol 1.4 % (CHLORASEPTIC) spray - for patients who cannot tolerate lozenges | 2 spray, Mouth/Throat, every 3 hours PRN, sore throat, Post-op |

Muscle Relaxants (Single Response)

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|-----|---|--|
| () | methocarbamol (ROBAXIN) 500 mg in sodium chloride 0.9 % 100 mL IVPB | 500 mg, intravenous, for 60 Minutes, every 8 hours PRN, muscle spasms, Post-op |
| () | methocarbamol (ROBAXIN) tablet | 500 mg, oral, every 8 hours PRN, muscle spasms, Post-op |
| () | cyclobenzaprine (FLEXERIL) tablet | 5 mg, oral, every 8 hours PRN, muscle spasms, Post-op |

Muscle Relaxants - Refractory Treatments (Single Response)

| | |
|--|---|
| <input type="checkbox"/> diazepam (VALIUM) injection | 2.5 mg, intravenous, every 8 hours PRN, muscle spasms, inadequate muscle spasm relief following administration of other agents, Post-op Indication(s): Other Specify: Muscle Relaxant |
| <input type="checkbox"/> diazepam (VALIUM) tablet | 2.5 mg, oral, every 8 hours PRN, muscle spasms, inadequate muscle spasm relief following administration of other agents, Post-op Indication(s): Other Specify: Muscle Relaxant |

Antiemetics

| | |
|---|--|
| <input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral (Selection Required) | "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 type="checkbox"/> promethazine (PHENERGAN) IV or Oral or Rectal | "Or" Linked Panel |
| <input 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 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 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. |
| <input type="checkbox"/> scopolamine (TRANSDERM-SCOP) 1.5 mg (1 mg over 3 days) - For Patients LESS than 65 years old | 1 patch, transdermal, for 72 Hours, every 72 hours, Post-op |

PRN Medications - Symptom Management

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|---|---|
| <input checked="" type="checkbox"/> acetaminophen (TYLENOL) tablet | 650 mg, oral, every 6 hours PRN, fever, Temperature greater than 101 F, Post-op |
| <input type="checkbox"/> Itching - Neurosurgery medications (Single Response) | Avoid diphenhydramine use in patients over 70 years old when possible. |
| <input type="checkbox"/> cetirizine (Zyrtec) tablet | 5 mg, oral, daily PRN, itching, Post-op |
| <input type="checkbox"/> diphenhydrAMINE (BENADRYL) injection | 12.5 mg, intravenous, every 12 hours PRN, itching, Post-op |

PRN Medications - Bowel Management

| | |
|--|---|
| <input checked="" type="checkbox"/> polyethylene glycol (MIRALAX) packet 17 gram | 17 g, oral, 2 times daily, Post-op |
| <input checked="" type="checkbox"/> docusate sodium (COLACE) capsule | 100 mg, oral, 2 times daily, Post-op |
| <input type="checkbox"/> magnesium hydroxide suspension | 30 mL, oral, daily PRN, constipation, Post-op |
| <input type="checkbox"/> bisacodyl (DULCOLAX) EC tablet | 5 mg, oral, daily PRN, constipation, Post-op |
| <input type="checkbox"/> bisacodyl (DULCOLAX) suppository | 10 mg, rectal, daily PRN, constipation, Post-op |
| <input type="checkbox"/> magnesium citrate solution | 150 mL, oral, daily PRN, constipation, For 2 Doses, Post-op |

PRN Medications - Bowel Management

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|---|-------------------------------|
| <input type="checkbox"/> saline, mineral oil, glycerin (S.M.O.G.) enema | 180 mL, rectal, once, Post-op |
|---|-------------------------------|

PRN Medications - Pain - Pain Score (1-3) (Single Response)

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|---|--|
| <input type="checkbox"/> traMADol (ULTRAM) tablet | 25 mg, oral, every 4 hours PRN, mild pain (score 1-3), Post-op Maximum Daily Dose: 200 mg/day |
| <input type="checkbox"/> traMADoL (ULTRAM) tablet | 50 mg, oral, every 6 hours PRN, mild pain (score 1-3), Post-op Maximum Daily Dose: 200 mg/day |

PCA Medications (Single Response)

| | |
|---|---|
| <input type="checkbox"/> morPHINE PCA 30 mg/30 mL | |
| <input type="checkbox"/> morPHINE 30 mg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 1 mg Lockout Interval: Not Ordered Basal Rate: 0 mg/hr MAX (Four hour dose limit): 20 mg intravenous, continuous, Post-op Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patients ages 19-59 years old with normal renal function, may bolus {Bolus Dose:26657::"2"} mg every {Bolus Frequency:26659::"3"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26660::"0.5"} mg ONCE. Adjust doses for age, renal function or other factors. |
| <input type="checkbox"/> Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change |
| <input type="checkbox"/> Pasero Opioid-induced Sedation Scale | Routine, Once |
| <input type="checkbox"/> Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy |
| <input type="checkbox"/> Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention |
| <input type="checkbox"/> naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3)., Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |
| <input type="checkbox"/> hydromorPHONE PCA (DILAUDID) 15 mg/30 mL | |
| <input type="checkbox"/> hydromorPHONE (DILAUDID) 15 mg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 0.2 mg Lockout: Not Ordered Basal Rate: 0 mg/hr MAX (Four hour dose limit): 3 mg intravenous, continuous, Post-op Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patients ages 19-59 years old with normal renal function, may bolus {Bolus Dose:26662::"0.2"} mg every {Bolus Frequency:26663::"3"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26664::"0.1"} mg ONCE. Adjust doses for age, renal function or other factors. Turn Off PCA Continuous Dose (Basal Rate) On Date: Turn Off PCA Continuous Dose (Basal Rate) At Time: |

| | |
|--|--|
| [] Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change |
| [] Pasero Opioid-induced Sedation Scale | Routine, Once |
| [] Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy |
| [] Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention |
| [] naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3)., Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |
| () fentaNYL PCA (SUBLIMAZE) 1500 mcg/30 mL | |
| [] fentaNYL (SUBLIMAZE) 1500 mcg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 10 mcg Lockout: Not Ordered Basal Rate: 0 mcg/hr Four Hour Dose Limit: 150 mcg intravenous, continuous, Post-op **Due to fentaNYL 600 mcg/30 mL shortages, the new standard for all facilities will be fentaNYL 1500 mcg/30 mL. This concentration is 2.5 x more concentrated.** Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patient 19-59 years old, may bolus {Bolus Dose:26656::"25"} mcg every {Bolus Frequency:26655::"2"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26654::"10"} mcg ONCE. Adjust doses for age, renal function or other factors. Turn Off PCA Continuous Dose (Basal Rate) On Date: Turn Off PCA Continuous Dose (Basal Rate) At Time: |
| [] Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change |
| [] Pasero Opioid-induced Sedation Scale | Routine, Once |
| [] Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy |

| | |
|---|---|
| <input type="checkbox"/> Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention |
| <input type="checkbox"/> naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3)., Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |

PCA Medications - HMSL, HMW, HMSTC, HMSTJ Only (Single Response)

() morPHINE PCA 30 mg/30 mL

| | |
|---|---|
| <input type="checkbox"/> morPHINE 30 mg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 1 mg Lockout Interval: Not Ordered Basal Rate: 0 mg/hr MAX (Four hour dose limit): 20 mg intravenous, continuous, Post-op Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patients ages 19-59 years old with normal renal function, may bolus {Bolus Dose:26657::"2"} mg every {Bolus Frequency:26659::"3"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26660::"0.5"} mg ONCE. Adjust doses for age, renal function or other factors. |
| <input type="checkbox"/> Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change, Post-op |
| <input type="checkbox"/> Pasero Opioid-induced Sedation Scale | Routine, Once |
| <input type="checkbox"/> Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy, Post-op |
| <input type="checkbox"/> Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention, Post-op |
| <input type="checkbox"/> naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3)., Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |

() hydromorPHONE PCA (DILAUDID) 15 mg/30 mL

| | |
|---|---|
| <input type="checkbox"/> hydromorPHONE (DILAUDID) 15 mg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 0.2 mg Lockout: Not Ordered Basal Rate: 0 mg/hr MAX (Four hour dose limit): 3 mg intravenous, continuous, Post-op Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patients ages 19-59 years old with normal renal function, may bolus {Bolus Dose:26662::"0.2"} mg every {Bolus Frequency:26663::"3"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26664::"0.1"} mg ONCE. Adjust doses for age, renal function or other factors. Turn Off PCA Continuous Dose (Basal Rate) On Date: Turn Off PCA Continuous Dose (Basal Rate) At Time: |
| <input type="checkbox"/> Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change, Post-op |
| <input type="checkbox"/> Pasero Opioid-induced Sedation Scale | Routine, Once |
| <input type="checkbox"/> Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy, Post-op |
| <input type="checkbox"/> Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention, Post-op |
| <input type="checkbox"/> naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3)., Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |
| () fentaNYL PCA (SUBLIMAZE) 600 mcg/30 mL | |
| <input type="checkbox"/> fentaNYL (SUBLIMAZE) 600 mcg/30 mL PCA | Nurse Loading Dose: Not Ordered PCA Dose: 10 mcg Lockout Interval: Not Ordered Basal Rate: 0 mcg/hr MAX (Four hour dose limit): 150 mcg intravenous, continuous, Post-op Management of breakthrough pain. Administer only if respiratory rate 12 per minute or more and POSS level of 2 or less. If more than 2 bolus doses in 12 hours or if pain persists after increase in demand dose, call ordering prescriber. For breakthrough pain in patient 19-59 years old, may bolus {Bolus Dose:26656::"25"} mcg every {Bolus Frequency:26655::"2"} hours as needed. If pain persists, may increase PCA demand dose by {PCA Dose:26654::"10"} mcg ONCE. Adjust doses for age, renal function or other factors. Turn Off PCA Continuous Dose (Basal Rate) On Date: Turn Off PCA Continuous Dose (Basal Rate) At Time: |

| | |
|---|--|
| <input type="checkbox"/> Vital signs - T/P/R/BP | Routine, Per unit protocol - Initially and every 30 minutes for 1 hour after PCA started, bolus administration or dose change; then - Every hour x 2 starting second hour after PCA started, bolus administered or dose change; then - Every 4 hours until PCA therapy is discontinued. - Immediately following PCA administration tubing change, Post-op |
| <input type="checkbox"/> Richmond agitation sedation scale | Routine, Once Hold infusion daily at: Target RASS: BIS Monitoring (Target BIS: 40-60): 60 minutes after administration of pain medication AND every 4 hours. Assess and document side effects of at least every 4 hours for duration of therapy and when patient complains of pain and/or side effects., Post-op |
| <input type="checkbox"/> Notify Physician (Specify) | Routine, Until discontinued, Starting S, - PCA pump infusion discontinued for any reason - Inadequate analgesia - Prior to administration of any other narcotics, antiemetics, or sedatives other than those ordered by the prescriber responsible for IV PCA therapy - PCA pump discontinued by any service other than the prescriber responsible for IV PCA therapy, Post-op |
| <input type="checkbox"/> Stop the PCA pump and call ordering physician and/or CERT team for any of the following: | Routine, Until discontinued, Starting S, - Respiratory rate 10 per minute or less - Severe and/or recent confusion or disorientation - POSS sedation level 4: Somnolent and difficult to arouse - Sustained hypotension (SBP less than 90) - Excessive nausea or vomiting - Urinary retention, Post-op |
| <input type="checkbox"/> naloxone (NARCAN) 0.4 mg/mL injection 0.2 mg | 0.2 mg, intravenous, once PRN, respiratory depression, as needed for respiratory rate 8 per minute or less OR patient somnolent and difficult to arouse (POSS GREATER than 3), Post-op Repeat Naloxone 0.2 mg once in 2 minutes if necessary (MAXIMUM 0.4 mg). If naloxone is needed, please call the ordering physician and/or CERT team. Monitor vital signs (pulse oximetry, P/R/BP) every 15 minutes for 3 times. |

PRN Medications - Pain - Pain Score (4-6) (Single Response)

| | |
|--|--|
| <input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet | 1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op |
| <input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) 300-30 mg per tablet | 1 tablet, oral, every 4 hours PRN, moderate pain (score 4-6), Post-op The use of codeine-containing products is contraindicated in patients LESS THAN 12 years of age. Is this patient OVER 12 years of age? Y/N: |
| <input type="checkbox"/> traMADol (ULTRAM) tablet | 50 mg, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum Daily Dose: 200 mg/day |
| <input type="checkbox"/> traMADoL (ULTRAM) tablet | 100 mg, oral, every 6 hours PRN, moderate pain (score 4-6) |

PRN Medications - Pain - Pain Score (7-10) (Single Response)

| | |
|--|---|
| <input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) 300-30 mg per tablet | 2 tablet, oral, every 6 hours PRN, severe pain (score 7-10), Post-op The use of codeine-containing products is contraindicated in patients LESS THAN 12 years of age. Is this patient OVER 12 years of age? Y/N: |
| <input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet | 2 tablet, oral, every 6 hours PRN, severe pain (score 7-10), Post-op |
| <input type="checkbox"/> traMADoL (ULTRAM) tablet | 50 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Maximum Daily Dose: 200 mg/day |

Breakthrough Pain (Single Response)

| | |
|---|--|
| <input type="checkbox"/> fentaNYL (SUBLIMAZE) injection | 25 mcg, intravenous, every 2 hour PRN, other, pain (score: 4-10) - inadequate pain relief following administration of oral agents, Post-op Administer after pain re-assessment for inadequate pain relief. |
| <input type="checkbox"/> morphine 2 mg/mL injection | 2 mg, intravenous, every 3 hours PRN, other, pain (score: 4-10) - inadequate pain relief following administration of oral agents, Post-op Administer after pain re-assessment for inadequate pain relief. |
| <input type="checkbox"/> HYDROmorphone (DILAUDID) injection | 0.5 mg, intravenous, every 3 hours PRN, other, pain (score: 4-10) - inadequate pain relief following administration of oral agents, Post-op Administer after pain re-assessment for inadequate pain relief. |

VTE

DVT Risk and Prophylaxis Tool (Single Response) (Selection Required)

VTE/DVT Risk Definitions

URL:

"\\appt1\epicapprod\Restricted\OrderSets\VTE DVTRISK DEFINITIONS.pdf"

[Anticoagulation Guide for COVID patients](#)

URL:

"https://formweb.com/files/houstonmethodist/documents/COVID-19 Anticoagulation Guideline - 8.20.2021v15.pdf"

| | |
|---|--|
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis with Risk Stratification (Single Response) (Selection Required) | |
| <input type="checkbox"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |

| | | |
|---|---|--|
| <input type="checkbox"/> | Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | | |
| <input type="checkbox"/> | High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> | Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> LOW Risk of DVT (Selection Required) | | |
| Low Risk Definition Age less than 60 years and NO other VTE risk factors | | |
| <input type="checkbox"/> Low Risk (Single Response) (Selection Required) | | |
| <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 (Selection Required) | | |
| Moderate Risk Definition Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. One or more of the following medical conditions: 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 Age 60 and above Central line History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hours Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission | | |
| <input type="checkbox"/> Moderate Risk (Selection Required) | | |
| <input type="checkbox"/> | Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required) | | |
| <input type="checkbox"/> | Contraindications exist for pharmacologic prophylaxis BUT order Sequential compression device | "And" Linked Panel |

| | | |
|--------------------------|---|--|
| <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"/> | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> | Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis | "And" Linked Panel |
| <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"/> | Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection (Single Response) (Selection Required) | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <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"/> | HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> | warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> | MODERATE Risk of DVT - Non-Surgical (Selection Required) | |

Moderate Risk Definition

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

One or more of the following medical conditions:

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

Age 60 and above

Central line

History of DVT or family history of VTE

Anticipated length of stay GREATER than 48 hours

Less than fully and independently ambulatory

Estrogen therapy

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Moderate Risk (Selection Required)

Moderate risk of VTE Routine, Once, PACU & Post-op

Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)

Contraindications exist for pharmacologic prophylaxis - **"And" Linked Panel**
Order Sequential compression device

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following
contraindication(s):
PACU & Post-op

Place/Maintain sequential compression device continuous Routine, Continuous, PACU & Post-op

Contraindications exist for pharmacologic prophylaxis **"And" Linked Panel**
AND mechanical prophylaxis

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following
contraindication(s):
PACU & Post-op

Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following
contraindication(s):
PACU & Post-op

enoxaparin (LOVENOX) injection (Single Response) (Selection Required)

enoxaparin (LOVENOX) syringe 40 mg, subcutaneous, daily at 1700, Starting S+1, PACU & Post-op
Indication(s): VTE Prophylaxis

patients with CrCL LESS than 30 mL/min 30 mg, subcutaneous, daily at 1700, Starting S+1, PACU & Post-op
For Patients with CrCL LESS than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min 30 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1, PACU & Post-op
For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min 40 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1, PACU & Post-op
For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min
Indication(s): VTE Prophylaxis

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"/> | HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> | warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | | |
| <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"/> HIGH Risk of DVT - Surgical (Selection Required) | | |
| High Risk Definition | | |
| Both pharmacologic AND mechanical prophylaxis must be addressed. | | |
| One or more of the following medical conditions: | | |
| Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) | | |
| Severe fracture of hip, pelvis or leg | | |
| Acute spinal cord injury with paresis | | |
| Multiple major traumas | | |
| Abdominal or pelvic surgery for CANCER | | |
| Acute ischemic stroke | | |
| History of PE | | |
| <input type="checkbox"/> High Risk (Selection Required) | | |
| <input type="checkbox"/> | High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required) | | |
| <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 for VTE Prophylaxis (Single Response) | | |
| <input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Daily at 1700 | | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 30 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Every 12 Hours | | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 30 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Daily at 1700 | | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 40 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Every 12 Hours | | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 40 mg, subcutaneous, every 12 hours Indication(s): |
| <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"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, Starting S+1, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> HIGH Risk of DVT - Non-Surgical (Selection Required) | |
| High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE | |
| <input type="checkbox"/> High Risk (Selection Required) | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required) | |
| <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) (Selection Required) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700, Starting S, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700, Starting S, PACU & Post-op For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <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"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> HIGH Risk of DVT - Surgical (Hip/Knee) (Selection Required) | |
| High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE | |
| <input type="checkbox"/> High Risk (Selection Required) | |
| <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) (Selection Required) | |
| <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"/> 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"/> Apixaban and Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> apixaban (ELIQUIS) tablet | 2.5 mg, oral, 2 times daily, Starting S+1, PACU & Post-op Indications: VTE prophylaxis |
| <input type="checkbox"/> Pharmacy consult to monitor apixaban (ELIQUIS) therapy | STAT, Until discontinued, Starting S Indications: VTE prophylaxis |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op For Patients with CrCL LESS than 30 mL/min. Indication(s): VTE Prophylaxis |
| <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, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. Indication(s): VTE Prophylaxis |

| | |
|--|--|
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 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, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <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"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> Rivaroxaban and Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (TIME CRITICAL), PACU & Post-op Indications: VTE prophylaxis |
| <input type="checkbox"/> Pharmacy consult to monitor rivaroxaban (XARELTO) therapy | STAT, Until discontinued, Starting S Indications: VTE prophylaxis |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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 |

DVT Risk and Prophylaxis Tool (Single Response)

VTE/DVT Risk Definitions

URL:

"\\appt1\epicapprod\Restricted\OrderSets\VTE\DVTRISK DEFINITIONS.pdf"

URL:

"https://formweb.com/files/houstonmethodist/documents/COVID-19 Anticoagulation Guideline - 8.20.2021v15.pdf"

| | |
|---|--|
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis with Risk Stratification (Single Response) (Selection Required) | |
| <input type="checkbox"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required) | |
| <input type="checkbox"/> High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis | 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"/> Place sequential compression device (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"/> LOW Risk of DVT (Selection Required) | |
| Low Risk Definition Age less than 60 years and NO other VTE risk factors | |
| <input type="checkbox"/> Low Risk (Single Response) (Selection Required) | |

| | |
|--|---|
| <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 |
|--|---|

MODERATE Risk of DVT - Surgical (Selection Required)

Moderate Risk Definition

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

One or more of the following medical conditions:

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

Age 60 and above

Central line

History of DVT or family history of VTE

Anticipated length of stay GREATER than 48 hours

Less than fully and independently ambulatory

Estrogen therapy

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Moderate Risk (Selection Required)

| | |
|---|-------------------------------|
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
|---|-------------------------------|

Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required)

| | |
|--|---------------------------|
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis BUT order Sequential compression device | "And" Linked Panel |
|--|---------------------------|

| | |
|--|---|
| <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"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|--|-------------------------------------|

| | |
|---|---------------------------|
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis | "And" Linked Panel |
|---|---------------------------|

| | |
|--|---|
| <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"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
|---|--|

enoxaparin (LOVENOX) injection (Single Response) (Selection Required)

| | |
|---|--|
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
|---|--|

| | |
|---|--|
| <input type="checkbox"/> patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis |
|---|--|

| | |
|---|--|
| <input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
|---|--|

| | |
|--|---|
| <input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
|--|---|

| | |
|---|--|
| <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"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> MODERATE Risk of DVT - Non-Surgical (Selection Required) | |
| Moderate Risk Definition Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. One or more of the following medical conditions: 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 Age 60 and above Central line History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hours Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission | |
| <input type="checkbox"/> Moderate Risk (Selection Required) | |
| <input type="checkbox"/> Moderate risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required) | |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis - Order Sequential compression device | "And" Linked Panel |
| <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"/> Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| <input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis | "And" Linked Panel |

| | |
|--|--|
| <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"/> Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| <input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required) | |
| <input type="checkbox"/> enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700, Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700, Starting S+1, PACU & Post-op For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <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"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> HIGH Risk of DVT - Surgical (Selection Required) | |
| High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE | |
| <input type="checkbox"/> High Risk (Selection Required) | |

| | | |
|--------------------------|--|---|
| <input type="checkbox"/> | High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> | High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required) | |
| <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 for VTE Prophylaxis (Single Response) | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) 30 mg Daily at 1700 | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 30 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> | enoxaparin (LOVENOX) 30 mg Every 12 Hours | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 30 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> | enoxaparin (LOVENOX) 40 mg Daily at 1700 | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 40 mg, subcutaneous, daily at 1700 Indication(s): |
| <input type="checkbox"/> | enoxaparin (LOVENOX) 40 mg Every 12 Hours | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) injection | 40 mg, subcutaneous, every 12 hours Indication(s): |
| <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"/> | HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, Starting S+1, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> | warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> | HIGH Risk of DVT - Non-Surgical (Selection Required) | |
| | High Risk Definition | |
| | Both pharmacologic AND mechanical prophylaxis must be addressed. | |
| | One or more of the following medical conditions: | |
| | Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) | |
| | Severe fracture of hip, pelvis or leg | |
| | Acute spinal cord injury with paresis | |
| | Multiple major traumas | |
| | Abdominal or pelvic surgery for CANCER | |
| | Acute ischemic stroke | |
| | History of PE | |
| <input type="checkbox"/> | High Risk (Selection Required) | |

| | | |
|--------------------------|--|--|
| <input type="checkbox"/> | High risk of VTE | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> | High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required) | |
| <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) (Selection Required) | |
| <input type="checkbox"/> | enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700, Starting S, PACU & Post-op Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700, Starting S, PACU & Post-op For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <input type="checkbox"/> | patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| <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"/> | HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, PACU & Post-op For patients with weight GREATER than 100 kg. |
| <input type="checkbox"/> | warfarin (COUMADIN) tablet | oral, daily at 1700, 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) (Selection Required) | |
| <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"/> | HIGH Risk of DVT - Surgical (Hip/Knee) (Selection Required) | |
| | High Risk Definition | |
| | Both pharmacologic AND mechanical prophylaxis must be addressed. | |
| | One or more of the following medical conditions: | |
| | Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) | |
| | Severe fracture of hip, pelvis or leg | |
| | Acute spinal cord injury with paresis | |
| | Multiple major traumas | |
| | Abdominal or pelvic surgery for CANCER | |
| | Acute ischemic stroke | |
| | History of PE | |
| <input type="checkbox"/> | High Risk (Selection Required) | |

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|--------------------------|--|--|
| <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) (Selection Required) | |
| () | Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () | aspirin chewable tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| () | aspirin (ECOTRIN) enteric coated tablet | 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| () | Apixaban and Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> | apixaban (ELIQUIS) tablet | 2.5 mg, oral, 2 times daily, Starting S+1, PACU & Post-op Indications: VTE prophylaxis |
| <input type="checkbox"/> | Pharmacy consult to monitor apixaban (ELIQUIS) therapy | STAT, Until discontinued, Starting S Indications: VTE prophylaxis |
| () | enoxaparin (LOVENOX) injection (Single Response) (Selection Required) | |
| () | enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| () | enoxaparin (LOVENOX) syringe | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1, PACU & Post-op Indication(s): VTE Prophylaxis |
| () | enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600, Starting S+1, PACU & Post-op For Patients with CrCL LESS than 30 mL/min. Indication(s): VTE Prophylaxis |
| () | 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, PACU & Post-op For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. Indication(s): VTE Prophylaxis |
| () | enoxaparin (LOVENOX) syringe - For Patients weight between 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, PACU & Post-op For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis |
| () | 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): |
| () | heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op |
| () | 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. |
| () | HEParin (porcine) injection - For Patients with weight GREATER than 100 kg | 7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op For patients with weight GREATER than 100 kg. |
| () | Rivaroxaban and Pharmacy Consult (Selection Required) | |
| <input type="checkbox"/> | rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (TIME CRITICAL), PACU & Post-op Indications: VTE prophylaxis |
| <input type="checkbox"/> | Pharmacy consult to monitor rivaroxaban (XARELTO) therapy | STAT, Until discontinued, Starting S Indications: VTE prophylaxis |
| () | warfarin (COUMADIN) tablet | oral, daily at 1700, Starting S+1, PACU & Post-op Indication: |
| () | Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |

Mechanical Prophylaxis (Single Response) (Selection Required)

- | | |
|--|--|
| <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 |

Labs

Laboratory

Type and screen

Type and screen Once, PACU & Post-op

ABO and Rh confirmation Once, Blood Bank Confirmation

Hemoglobin and hematocrit Once
In Recovery room., PACU & Post-op

Basic metabolic panel Once, PACU & Post-op

CBC with platelet and differential Once, PACU & Post-op

Partial thromboplastin time Once, PACU & Post-op

Prothrombin time with INR Once, PACU & Post-op

Calcium level Once, PACU & Post-op

Magnesium level Once, PACU & Post-op

Phosphorus level Once, PACU & Post-op

Blood gas, arterial Once, PACU & Post-op

Urinalysis screen and microscopy, with reflex to culture Once
Specimen Source: Urine
Specimen Site:
PACU & Post-op

Labs - AM

Basic metabolic panel AM draw, Starting S+1 For 1 Occurrences, PACU & Post-op

CBC with platelet and differential AM draw, Starting S+1 For 1 Occurrences, PACU & Post-op

Partial thromboplastin time AM draw, Starting S+1 For 1 Occurrences, PACU & Post-op

Prothrombin time with INR AM draw, Starting S+1 For 1 Occurrences, PACU & Post-op

Labs - AM Daily x 3

Hemoglobin AM draw repeats For 3 Occurrences, PACU & Post-op

Imaging

CT

CT Cervical Spine Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1 , PACU & Post-op

CT Thoracic Spine Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1 , PACU & Post-op

X-ray

Chest 1 Vw Portable Routine, 1 time imaging, Starting S at 1:00 AM For 1 , PACU & Post-op

Chest 1 Vw Portable in AM Routine, 1 time imaging, Starting S+1 For 1 , PACU & Post-op

XR Spine Scoliosis 2-3 Views Routine, 1 time imaging, Starting S at 1:00 AM For 1
Please add 32 millimeter image calibration necklace to the field of view. AP and Lateral view that includes C2 and femoral heads in single shot with patient standing with hips and knees extended., PACU & Post-op

Cervical Spine 2 Or 3 Vw Routine, 1 time imaging, Starting S at 1:00 AM For 1 , PACU & Post-op

Respiratory

Respiratory

| | |
|---|---|
| <input checked="" type="checkbox"/> Oxygen therapy - Simple face mask | Routine, Continuous Device: Simple Face Mask Rate in liters per minute: 6 Lpm Rate in tenths of a liter per minute: O2 %: Titrate to keep O2 Sat Above: 92% Indications for O2 therapy: Immediate post-op period Device 2: Device 3: Wean prn., PACU & Post-op |
| <input type="checkbox"/> Incentive spirometry | Routine, Once, PACU & Post-op |
| <input type="checkbox"/> Mechanical ventilation | Routine, PACU & Post-op Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: |

Consults

For Physician Consult orders use sidebar

Ancillary Consults

| | |
|---|---|
| <input type="checkbox"/> Consult to Case Management | Consult Reason: PACU & Post-op |
| <input type="checkbox"/> Consult to Social Work | Reason for Consult: PACU & Post-op |
| <input checked="" type="checkbox"/> Consult PT eval and treat | Reasons for referral to Physical Therapy (mark all applicable): Post Neuromuscular or Musculoskeletal Surgery Care. Are there any restrictions for positioning or mobility? Please provide safe ranges for HR, BP, O2 saturation(if values are very abnormal): Weight Bearing Status: PACU & Post-op |
| <input type="checkbox"/> Consult PT wound care | Special Instructions: Location of Wound? PACU & Post-op |
| <input checked="" type="checkbox"/> Consult OT eval and treat | Reason for referral to Occupational Therapy (mark all that apply): Decline in Activities of Daily Living performance from baseline (bathing, dressing, toileting, grooming) Are there any restrictions for positioning or mobility? Please provide safe ranges for HR, BP, O2 saturation(if values are very abnormal): Weight Bearing Status: PACU & Post-op |
| <input type="checkbox"/> Consult to Nutrition Services | Reason For Consult? Purpose/Topic: PACU & Post-op |
| <input type="checkbox"/> Consult to Spiritual Care | Reason for consult? PACU & Post-op |
| <input type="checkbox"/> Consult to Speech Language Pathology | Routine, Once Reason for consult: PACU & Post-op |
| <input type="checkbox"/> Consult to Wound Ostomy Care nurse | Reason for consult: Reason for consult: Reason for consult: Reason for consult: Consult for NPWT: Reason for consult: Reason for consult: PACU & Post-op |

| | |
|--|--|
| <input type="checkbox"/> Consult to Respiratory Therapy | Reason for Consult? Post neuromuscular or musculoskeletal surgery care PACU & Post-op |
| Physician Consults | |
| <input checked="" type="checkbox"/> Consult Intensive Care | Reason for Consult? Decline in ADL performance from baseline Patient/Clinical information communicated? Telephone Patient/clinical information communicated? Telephone PACU & Post-op |
| <input type="checkbox"/> Consult Physical Medicine Rehab | Reason for Consult? Patient/Clinical information communicated? Patient/clinical information communicated? PACU & Post-op |