Mini-Maze Post-Op ICU [4113]

| General | |
|---|--|
| | |
| Common Present on Admission Diagnosis | |
| [] Acidosis | Post-op |
| [] Acute Post-Hemorrhagic Anemia | Post-op |
| [] Acute Renal Failure | Post-op |
| [] Acute Respiratory Failure | Post-op |
| [] Acute Thromboembolism of Deep Veins of Lower Extremities | Post-op |
| [] Anemia | Post-op |
| [] Bacteremia | Post-op |
| [] Bipolar disorder, unspecified | Post-op |
| [] Cardiac Arrest | Post-op |
| [] Cardiac Dysrhythmia | Post-op |
| [] Cardiogenic Shock | Post-op |
| [] Decubitus Ulcer | Post-op |
| Dementia in Conditions Classified Elsewhere | Post-op |
| [] Disorder of Liver | Post-op |
| [] Electrolyte and Fluid Disorder | Post-op |
| [] Intestinal Infection due to Clostridium Difficile | Post-op |
| [] Methicillin Resistant Staphylococcus Aureus Infection | Post-op |
| [] Obstructive Chronic Bronchitis with Exacerbation | Post-op |
| [] Other Alteration of Consciousness | Post-op |
| [] Other and Unspecified Coagulation Defects | Post-op |
| [] Other Pulmonary Embolism and Infarction | Post-op |
| [] Phlebitis and Thrombophlebitis | Post-op |
| [] Protein-calorie Malnutrition | Post-op |
| [] Psychosis, unspecified psychosis type | Post-op |
| [] Schizophrenia Disorder | Post-op |
| [] Sepsis | Post-op |
| [] Septic Shock | Post-op |
| [] Septicemia | Post-op |
| [] Type II or Unspecified Type Diabetes Mellitus with Mention of Complication, Not Stated as Uncontrolled | Post-op |
| [] Urinary Tract Infection, Site Not Specified | Post-op |
| Elective Outpatient, Observation, or Admission (Single | Response) |
| () Elective outpatient procedure: Discharge following | Routine, Continuous, PACU & Post-op |
| routine recovery | |
| () Outpatient observation services under general | Diagnosis: |
| supervision | Admitting Physician: Patient Condition: |
| | Patient Condition: Bed request comments: |
| | PACU & Post-op |
| () Outpatient in a bed - extended recovery | Diagnosis: |
| () Outpution in a bod Extended recovery | Admitting Physician: |
| | Bed request comments: |
| | PACU & Post-op |
| () Admit to Inpatient | Diagnosis: |
| () | Admitting Physician: |
| | Level of Care: |
| | Patient Condition: |
| | Bed request comments: |
| | Certification: I certify that based on my best clinical judgment |
| | and the patient's condition as documented in the HP and |
| | progress notes, I expect that the patient will need hospital |
| | services for two or more midnights. |
| <u> </u> | PACU & Post-op |
| Printed on 9/17/2019 at 2:30 PM from SUP | Page 1 of 2 |

Admission or Observation (Single Response) Patient has active outpatient status order on file

| () Admit to Inpatient | Diagnosis: |
|--|---|
| | Admitting Physician: |
| | Level of Care: Patient Condition: |
| | |
| | Bed request comments: |
| | Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and |
| | progress notes, I expect that the patient will need hospital |
| | services for two or more midnights. PACU & Post-op |
|) Outpatient observation services under general | Diagnosis: |
| supervision | Admitting Physician: |
| · | Patient Condition: |
| | Bed request comments: |
| | PACU & Post-op |
|) Outpatient in a bed - extended recovery | Diagnosis: |
| | Admitting Physician: |
| | Bed request comments: |
| | PACU & Post-op |
|) Transfer patient | Level of Care: |
| | Bed request comments: |
| | Scheduling/ADT |
|) Return to previous bed | Routine, Until discontinued, Starting S, Scheduling/ADT |
| Admission (Single Response) Patient has active status order on file | |
| () Admit to inpatient | Diagnosis: |
| | Admitting Physician: Level of Care: |
| | Patient Condition: |
| | Bed request comments: |
| | Certification: I certify that based on my best clinical judgment |
| | and the patient's condition as documented in the HP and |
| | and the battern's condition as documented in the HF and |
| | • |
| | progress notes, I expect that the patient will need hospital |
| | • |
|) Transfer patient | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: |
|) Transfer patient | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: |
| | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT |
| | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: |
| | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT |
| () Return to previous bed Transfer (Single Response) Patient has active inpatient status order on file | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT |
| () Return to previous bed Fransfer (Single Response) Patient has active inpatient status order on file | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: |
| () Return to previous bed Transfer (Single Response) Patient has active inpatient status order on file | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: |
| () Return to previous bed Fransfer (Single Response) Patient has active inpatient status order on file () Transfer patient | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: |
| () Return to previous bed Fransfer (Single Response) Patient has active inpatient status order on file () Transfer patient () Return to previous bed | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: Scheduling/ADT |
| () Return to previous bed Transfer (Single Response) Patient has active inpatient status order on file () Transfer patient () Return to previous bed Code Status | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT |
| 7) Return to previous bed Fransfer (Single Response) Patient has active inpatient status order on file 7) Transfer patient 8) Return to previous bed | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: Scheduling/ADT |
| () Return to previous bed Transfer (Single Response) Patient has active inpatient status order on file () Transfer patient () Return to previous bed Code Status | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Code Status decision reached by: |
| () Return to previous bed Transfer (Single Response) Patient has active inpatient status order on file () Transfer patient () Return to previous bed Code Status [] Full Code | progress notes, I expect that the patient will need hospital services for two or more midnights. PACU & Post-op Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Level of Care: Bed request comments: Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Routine, Until discontinued, Starting S, Scheduling/ADT Code Status decision reached by: |

| [] Consult to Palliative Care Service | Priority: |
|--|--|
| | Reason for Consult? |
| | Order? |
| | Name of referring provider: |
| | Enter call back number: |
| [] Consult to Social Work | Reason for Consult: |
| | Post-op |
| [] Modified Code | Does patient have decision-making capacity? |
| | Modified Code restrictions: |
| | Post-op |
| [] Treatment Restrictions | Treatment Restriction decision reached by: |
| []aament toomenene | Specify Treatment Restrictions: |
| | Post-op |
| | |
| Isolation | |
| [1] Airborno inclotion status | |
| [] Airborne isolation status | D |
| [] Airborne isolation status | Details |
| [] Mycobacterium tuberculosis by PCR - If you suspect | Once, Sputum, Post-op |
| Tuberculosis, please order this test for rapid diagnostics. | |
| [] Contact isolation status | Details |
| [] Droplet isolation status | Details |
| [] Enteric isolation status | Details |
| | |
| Precautions | |
| [] Aspiration procautions | Post on |
| [] Aspiration precautions | Post-op |
| [] Fall precautions | Increased observation level needed: Post-op |
| | POSI-00 |
| [1] Later properties | |
| [] Latex precautions | Post-op |
| Latex precautions Seizure precautions | Post-op Increased observation level needed: |
| | Post-op |
| [] Seizure precautions | Post-op Increased observation level needed: |
| | Post-op Increased observation level needed: |
| [] Seizure precautions Nursing | Post-op Increased observation level needed: |
| Nursing Vital Signs | Post-op Increased observation level needed: Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP | Post-op Increased observation level needed: |
| Nursing Vital Signs | Post-op Increased observation level needed: Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall,with assistance Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance Post-op Routine, Daily, Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall,with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) Specify: 35 |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights [X] Head of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall,with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) Specify: 35 Post-op |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall,with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) Specify: 35 Post-op Routine, Every hour, Starting S |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights [X] Head of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP, Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed, Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall, with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) Specify: 35 Post-op Routine, Every hour, Starting S Assessment to Perform: Cranial Nerves, Glasgow Coma |
| Nursing Vital Signs [X] Vital signs - T/P/R/BP [X] Hemodynamic Monitoring Activity [X] Dangle at bedside [X] Out of bed [X] Ambulate Nursing [X] Daily weights [X] Head of bed | Post-op Increased observation level needed: Post-op Routine, Per unit protocol, Post-op Routine, Continuous Measure: Arterial Line MAP,Arterial Line BP Post-op Routine, Once Begin on POD 0, Post-op Routine, Until discontinued, Starting S Specify: Out of bed,Up in chair Additional modifier: for meals Chair x 3 daily, Post-op Routine, 4 times daily Specify: in hall,with assistance Post-op Routine, Daily, Post-op Routine, Until discontinued, Starting S Head of bed: other degrees (specify) Specify: 35 Post-op Routine, Every hour, Starting S |

| x 2 consecutive hours or less than 240mL per 12 hour shift | , |
|---|--|
| mililiters/hour [X] Notify Physician -For urine output LESS THAN 100 ml/hr | Routine, Until discontinued, Starting S, Post-op |
| [X] Notify Physician - for chest output greater than 200 | SpO2 less than: 90 Routine, Until discontinued, Starting S, Post-op |
| | Respiratory rate greater than: 30 Respiratory rate less than: |
| | Heart rate less than (BPM): 40 |
| | Heart rate greater than (BPM): 120 |
| | Diastolic BP less than: MAP less than: 55 |
| | Diastolic BP greater than: |
| | Systolic BP less than: 80 |
| | Systolic BP greater than: 180 |
| | Temperature greater than: 102.5 Temperature less than: 95 |
| [X] Notify Physician for vitals: | Routine, Until discontinued, Starting S |
| Notify | |
| Wires | Before transfer out of ICU; if not already discontinued. , Post-op |
| [] Discontinue Pacemaker Generator and Insulate Pacer | 1 or POD 2. , Post-op Routine, Conditional Frequency For 1 Occurrences |
| | Foley reason for not removing MUST be documented on POD |
| [X] Foley catheter - discontinue | Routine, Conditional Frequency For 1 Occurrences 1) Remove Foley cath POD 1 or POD 2; If unable to remove |
| E-1 = | Before transfer out of ICU; if arterial line not already discontinued, Post-op |
| Discontinue [X] Discontinue arterial line | Routine, Conditional Frequency For 1 Occurrences |
| Piacontinua | . oc. op |
| | Options: Post-op |
| | AV Interval (milliseconds): |
| | Sensitivity Setting (millivolts): |
| | Ventrical Setting (MA): |
| [א] ו מספווומעפו ספונווואס | Atrial Setting (MA): |
| [X] Pacemaker settings | blood glucose greater than 300 mg / dL, Post-op Routine, Until discontinued, Starting S |
| | drip; Notify physician for blood glucose less than 70 mg/dL OF |
| | hours then change to every 4 hours if not started on an insulin |
| | Every hour For Until specified Monitor every hour for first 6 |
| [X] Bedside glucose | Routine, Every hour For Until specified (Q1 hour x 6) ONLY IF HISTORY OF DIABETES Routine, |
| IVI Dedeide elvesse | hours, Post-op |
| [X] Oral care | Routine, 2 times daily Every 12 hours Per CVICU protocol. Toothbrush every 12 |
| | Chest tube site care daily and prn per protocol, Post-op |
| [X] Tube site care (chest tube) | Post-op Routine, Per unit protocol |
| | Level of suction: 20 cm H2O |
| [X] Chest tube to continuous suction | Clean with CHG cloths, Post-op Routine, Until discontinued, Starting S |
| | Orders: Maintain |
| [X] Foley catheter care | Routine, 2 times daily |
| | To achieve body temperature of 98.6 F, Post-op |
| [X] Apply warming blanket (bair hugger) | Site: epicardial pacing wire site, Post-op Routine, Once For 1 Occurrences |
| | Site: |
| [X] Site care | Routine, Per unit protocol |

| Diet | |
|--|--|
| [X] Diet - Regular | Diet effective now, Starting S |
| | Diet(s): Regular |
| | Advance Diet as Tolerated? Liquid Consistency: |
| | Fluid Restriction: |
| | Foods to Avoid: |
| | Post-op |
| Prune Juice or Prunes | Routine, Until discontinued, Starting S |
| | Give with breakfast daily starting post op day 2, Post-op |
| IV Fluids | |
| V Fluids (Single Response) | |
| (X) lactated Ringer's infusion | 75 mL/hr, intravenous, continuous, Post-op |
| Medications | |
| Pharmacy Consults for Heparin Management | |
| [] Pharmacy consult to manage Heparin LOW dose | STAT, Until discontinued, Starting S |
| protocol (ACS/Stroke/Afib) - initiation bolus and infusion | Heparin Indication: |
| withOUT titration boluses | Specify: |
| II. Diaman Oraș lita Maran II. a ' OTANDADD I | Monitoring: Anti-Xa |
| [] Pharmacy Consult to Manage Heparin STANDARD dose | STAT, Until discontinued, Starting S Heparin Indication: |
| protocol (DVT/PE) - initiation bolus and infusion with titration boluses | Specify: Give initial Bolus |
| illiation bolases | Monitoring: Anti-Xa |
| PostOp Antibiotics: For Patients LESS than or EQUAL to 1 | 20 kg (Single Response) |
| (X) ceFAZolin (ANCEF) IV - For Patients LESS than or | 2 g, intravenous, every 8 hours, For 2 Doses, Post-op |
| EQUAL to 120 kg | Reason for Therapy: Surgical Prophylaxis |
| () If Beta-Lactam Allergic - vancomycin (VANCOCIN) IV | 15 mg/kg, intravenous, once, For 1 Doses, Post-op Administer 12 hours after procedure |
| | Reason for Therapy: Surgical Prophylaxis |
| Post-Op Antibiotics: For Patients GREATER than 120 kg (S | Single Response) |
| (X) ceFAZolin (ANCEF) IV - For Patients GREATER than | 3 g, intravenous, every 8 hours, For 2 Doses, Post-op |
| 120 kg | Reason for Therapy: Surgical Prophylaxis |
| () If Beta-Lactam Allergic - vancomycin (VANCOCIN) IV | 15 mg/kg, intravenous, once, For 1 Doses, Post-op |
| | Administer 12 hours after procedure |
| | Reason for Therapy: Surgical Prophylaxis |
| notropes | |
| DOPamine (INTROPIN) infusion | 2-20 mcg/kg/min, intravenous, continuous, Post-op |
| | Titrate for cardiac index GREATER than 2.2 or Mean Arterial |
| | Pressure GREATER than 60. Recommendation is to titrate |
| | with 1-10 mcg/kg/min. Notify intensivist when titration requires |
| | GREATER than 5 mcg/kg/min. Wean to off when parameters |
| The EDINEDILE AND ENAMENDED AND THE STATE OF | are satisfied. Discontinue in Epic when off for 4 hours. |
| [] EPINEPHrine (ADRENALIN) in sodium chloride 0.9 % 250 mL infusion | 2-30 mcg/min, intravenous, continuous, Post-op Titrate for cardiac index GREATER than 2.2 or Mean Arterial |
| 200 IIIL IIIIUSIOII | Pressure GREATER than 60. Recommendation is to titrate |
| | with 1-10 mcg/min. Notify intensivist when titration requires |
| | greater than 5 mcg/min. Wean to off when parameters are |
| | satisfied. Discontinue in Epic when off for 4 hours. |
| DOButamine (DOBUTREX) infusion | 0.5-20 mcg/kg/min, intravenous, continuous, Post-op |
| | Titrate for cardiac index GREATER than 2.2. |
| | Recommendation is to titrate with 2-10 mcg/kg/min. Notify |
| | intensivist when titration requires GREATER than 5 |
| | mcg/kg/min. Wean to off when parameters are satisfied. |
| | Discontinue in Epic when off for 4 hours. |

| [] milrinone (PRIMACOR) infusion | 0.125-0.75 mcg/kg/min, intravenous, continuous, Post-op Titrate for cardiac index GREATER than 2.2. Recommendation is to titrate with 0.25-0.75 mcg/kg/min. Notify intensivist when titration requires GREATER than 0.5 mcg/kg/min. Wean to off when parameters are satisfied. |
|--|---|
| Pressors | Discontinue in Epic when off for 4 hours. |
| [] vasopressin (PITRESSIN) 0.4 Units/mL in sodium chloride 0.9 % 100 mL infusion | 0.01-0.04 Units/min, intravenous, continuous, Post-op Titrate for mean arterial pressure GREATER than 60. Recommendation is to titrate with 0.02 to 0.1 units/min. Notify intensivist when titration requires greater than 0.06 units/min. Wean to off when parameters are satisfied. Discontinue vasopressin order in Epic when off for 4 hours. |
| [] norepinephrine (LEVOPHED) infusion | 4-50 mcg/min, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure GREATER than 60. Recommendation is to titrate with 2 to 12 mcg/min. Notify intensivist when titration requires greater than 8 mcg/min. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| [] phenylephrine (NEO-SYNEPHRINE) in sodium chloride 0.9 % 250 mL infusion | 5-150 mcg/min, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure GREATER than 60. Recommendation is to titrate with 100-180 mcg/min. Notify intensivist when titration requires GREATER than 150 mcg/min. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| IV infusion - Antihypertensives (Single Response) | |
| () niCARdipine (CARDENE) IV infusion | 2.5-15 mg/hr, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure 70-80 mmHg. Recommendation is to titrate with 5 to 15 mg/hr. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| () clevidipine (CLEVIPREX) infusion | 1-32 mg/hr, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure 70-80 mmHg. Recommendation is to titrate with 2 to 32 mg/hr. Notify intensivist when titration requires GREATER than 16 mg/hr. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| () diltiazem (CARDIZEM) infusion | 2.5-10 mg/hr, intravenous, continuous, Post-op |
| () nitroglycerin infusion | 5-200 mcg/min, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure GREATER than 60. Recommendation is to titrate with 10 to 200 mcg/min. Notify intensivist when titration requires GREATER than 100 mcg/min. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| () esmolol (BREVIBLOC) infusion | 50-300 mcg/kg/min, intravenous, continuous, Post-op Titrate for Mean Arterial Pressure from 70-80 mmHg. Recommendation is to titrate with 50-300 mcg/kg/min. Notify intensivist when titration requires GREATER than 200 mcg/kg/min. Wean to off when parameters are satisfied. Discontinue order in Epic when off for 4 hours. |
| () labetalol infusion | 2 mg/min, intravenous, continuous, Post-op [labetalol]HOLD parameters for this order: [labetalol]Contact Physician if: |
| Other Medications | |
| [] colchicine tablet FOR DIABETIC ONLY | 0.6 mg, oral, daily, Post-op For prevention of atrial fibrillation post cardiac surgery. Call provider for diarrhea. |
| [X]_Prednisolone oral titrate | "Followed by" Linked Panel |
| [X] predniSONE (DELTASONE) tablet | 20 mg, oral, 2 times daily, For 3 Doses, Post-op |
| Printed on 9/17/2019 at 2:30 PM from SLIP | Page 6 of |

| [X] predniSONE (DELTASONE) tablet | 15 mg, oral, 2 times daily, For 3 Doses, Post-op |
|--|--|
| [X] predniSONE (DELTASONE) tablet | 10 mg, oral, 2 times daily, For 3 Doses, Post-op |
| [X] predniSONE (DELTASONE) tablet | 10 mg, oral, 2 times daily, For 3 Doses, Post-op |
| [X] furosemide (LASIX) Oral or IV (Single Response) | |
| () furosemide (LASIX) tablet | 40 mg, oral, daily, Post-op If unable to swallow oral tablets, discontinue and change to IV daily. |
| () furosemide (LASIX) IV | 40 mg, intravenous, daily, Post-op if unable to swallow oral tablets. |
| ACE Inhibitors (Single Response) | |
| () captopril (CAPOTEN) tablet | 25 mg, oral, 3 times daily, Post-op Consult MD before administering if urine output less than 0.5 mL/kg/hour and creatinine greater than 1.3. HOLD parameters for this order: Hold Parameters requested HOLD for: Other Please specify: 90 HOLD for Heart Rate LESS than: Contact Physician if: |
| () enalapril (VASOTEC) tablet | 2.5 mg, oral, 2 times daily, Post-op Consult MD before administering if urine output less than 0.5 mL/kg/hr and creatinine greater than 1.3. HOLD parameters for this order: Hold Parameters requested HOLD for: Other Please specify: 90 HOLD for Heart Rate LESS than: Contact Physician if: |
| () lisinopril (PRINIVIL,ZESTRIL) tablet | 5 mg, oral, daily, Post-op Consult MD before administering if urine output less than 0.5 mL/kg/hr and creatinine greater than 1.3. HOLD parameters for this order: Hold Parameters requested HOLD for: Other Please specify: 90 mmHg HOLD for Heart Rate LESS than: Contact Physician if: |
| amIODarone (CORDARONE) 24-hr Infusions HARD-Stop (Sin | igle Response) |
| () Loading Dose and Maintenance Infusion (Single Response) | |
| Select Standard or Double concentration | |
| () Standard [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by STANDARD concentration 24-hour Infusion for Atrial Fibrillation- NOT HMWB | "Followed by" Linked Panel |
| [] amIODarone (CORDArone) 150 mg BOLUS | 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion | 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours, Post-op |
| [] REDUCE rate for amIODarone (CORDArone) 450 mg/ 250 mL NS | 0.5 mg/min, intravenous, once, Starting H+6 Hours, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |

| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion - 2nd bag | 0.5 mg/min, intravenous, continuous, Starting H+8 Hours, For 16 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a central line or PICC line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. |
|--|---|
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by STANDARD concentration 24-hour Infusion for Atrial Fibrillation-HMWB ONLY | "Followed by" Linked Panel |
| [] amIODarone (CORDArone) 150 mg BOLUS | 150 mg, intravenous, once, Starting S, For 1 Doses Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion | 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours |
| [] REDUCE rate for amIODarone (CORDArone) infusion | 0.5 mg/min, intravenous, once, Starting H+6 Hours, For 1 Doses Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion - 2nd bag | 0.5 mg/min, intravenous, continuous, Starting H+8 Hours, For 16 Hours Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a central line or PICC |
| | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. |
| () Double | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. |
| () Double [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation | line if infusion duration is GREATER THAN 24 hours. Use |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. "Followed by" Linked Panel 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation [] amIODarone (CORDArone) 150 mg BOLUS | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. "Followed by" Linked Panel 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours, Post-op 0.5 mg/min, intravenous, continuous, Starting H+6 Hours, For 18 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation [] amIODarone (CORDArone) 150 mg BOLUS [] amIODarone (CORDArone) 900 mg/ 250 mL NS [] REDUCE rate for amIODarone (CORDArone) 900 mg/ | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. "Followed by" Linked Panel 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours, Post-op 0.5 mg/min, intravenous, continuous, Starting H+6 Hours, For 18 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation [] amIODarone (CORDArone) 150 mg BOLUS [] amIODarone (CORDArone) 900 mg/ 250 mL NS [] REDUCE rate for amIODarone (CORDArone) 900 mg/ 250 mL infusion | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. "Followed by" Linked Panel 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours, Post-op 0.5 mg/min, intravenous, continuous, Starting H+6 Hours, For 18 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire |
| [] CENTRAL Line Administration: amIODarone (CORDArone) 150 mg LOADING Dose followed by DOUBLE concentration 24-hour Infusion for Atrial Fibrillation [] amIODarone (CORDArone) 150 mg BOLUS [] amIODarone (CORDArone) 900 mg/ 250 mL NS [] REDUCE rate for amIODarone (CORDArone) 900 mg/ 250 mL infusion () Maintenance Infusion (Single Response) Select Standard or Double Concentration | line if infusion duration is GREATER THAN 24 hours. Use 0.2 Micron Filter Tubing for administration. "Followed by" Linked Panel 150 mg, intravenous, once, Starting S, For 1 Doses, Post-op Patients should be monitored for QTc prolongation. Use 0.2 Micron Filter Tubing for administration. 1 mg/min, intravenous, continuous, Starting H+10 Minutes, For 6 Hours, Post-op 0.5 mg/min, intravenous, continuous, Starting H+6 Hours, For 18 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire |

| REDUCE rate for amIODarone (CORDArone) 450 mg/ | 0.5 mg/min, intravenous, once, Starting H+6 Hours, For 1 |
|--|---|
| 250 mL NS | Doses, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion - 2nd bag | 0.5 mg/min, intravenous, continuous, Starting H+8 Hours, For 16 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |
| [] NO LOADING DOSE - Central Line Administration: amIODarone (CORDArone) STANDARD concentration 24-hour Infusion for Atrial Fibrillation - HMWB Only | "Followed by" Linked Panel |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion | 1 mg/min, intravenous, continuous, For 6 Hours |
| [] REDUCE rate for amIODarone (CORDArone) 360 mg/ 200 mL NS | 0.5 mg/min, intravenous, once, Starting H+6 Hours, For 1 Doses Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |
| [] amIODarone 1.8 mg/mL (STANDARD concentration) infusion - 2nd bag | 0.5 mg/min, intravenous, continuous, Starting H+8 Hours, For 16 Hours Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |
| () Double (Single Response) | |
| NO LOADING DOSE - Central Line Administration: amIODarone (CORDArone) Double Concentration 24-hour Infusion for Atrial Fibrillation | "Followed by" Linked Panel |
| [] amIODarone (CORDArone) 900 mg/ 250 mL NS | 1 mg/min, intravenous, continuous, For 6 Hours, Post-op Start with 1 mg/min for 6 hours. Decrease to 0.5 mg/min for 18 hours. Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Use 0.2 Micron Filter Tubing for administration. |
| [] REDUCE rate for amIODarone (CORDArone) 900 mg/ 250 mL NS | 0.5 mg/min, intravenous, continuous, Starting H+6 Hours, For 18 Hours, Post-op Patients should be monitored for QTc prolongation. Use in-line filter to help prevent Phlebitis. HOLD Infuson for heart rate LESS THAN 60 or pauses GREATER THAN 2.5 seconds. MUST be infused via a CENTRAL line or PICC line. Do not take down 1st infusion until entire content of bag is infused. |

amioDarone (PACErone) tablet

You MUST be sure the oral tablet order is set to start TOMORROW with the start time set to 24 hours AFTER the start time of the INITIAL infusion order above.

| [] amIODarone (PACERONE) tablet **** You MUST CHANGE the START DATE to TOMORROW and set the Start TIME to be 24 hours after the Start Time of the Infusion | oral, every 24 hours, Starting H+24 Hours amiodarone (Pacerone) tablets must start 24 hours after the start of the infusion order. |
|--|--|
| Antiplatelet Agents (Single Response) | |
| () Loading Dose Followed By Maintenance (Single Response) | |
| () clopidogrel (PLAVIX) 300 mg Loading Dose followed by 75 mg Maintenance Dose and aspirin EC 81 mg tablet | |
| [] clopidogrel (PLAVIX) Loading and Maintenance doses | "Followed by" Linked Panel |
| [] Loading Dose - clopidogrel (PLAVIX) tablet | 300 mg, oral, once, For 1 Doses, Post-op Loading Dose |
| [] Maintenance Dose - clopidogrel (PLAVIX) tablet | 75 mg, oral, daily, Starting S+1, Post-op Maintenance Dose |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| () ticagrelor (BRILINTA) 180 mg Loading Dose followed by 90 mg Maintenance Dose and aspirin EC 81 mg tablet | |
| [] ticagrelor (BRILANTA) Oral Loading and Maintenance Doses | "Followed by" Linked Panel |
| [] Loading Dose - ticagrelor (BRILINTA) tablet | 180 mg, oral, once, For 1 Doses, Post-op Loading Dose |
| [] Maintenance Dose - ticagrelor (BRILINTA) tablet | 90 mg, oral, 2 times daily, Starting H+12 Hours, Post-op Maintenance Dose |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| () prasugrel (EFFIENT) 60 mg Loading Dose followed by 10 mg Maintenance Dose and aspirin EC 81 mg tablet | |
| [] prasugrel (EFFIENT) Loading and Maintenance Doses | "Followed by" Linked Panel |
| Maintenance Dose Instructions: | |
| Lower the dose to 5 mg for high risk patients (age GRE | ATER than or EQUAL to 75 OR weight LESS than 60 kg) |
| [] Loading Dose - prasugrel (EFFIENT) tablet | 60 mg, oral, once, For 1 Doses, Post-op Loading Dose |
| [] Maintenance Dose - prasugrel (EFFIENT) tablet | 10 mg, oral, daily, Starting H+24 Hours, Post-op Maintenance Dose |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| [] ** DO NOT REMOVE ** Pharmacy Consult to educate | |
| patient on prasugrel (EFFIENT) | 0747 0 5 4 0 |
| [] Pharmacy Consult to educate patient on prasugrel (EFFIENT) | STAT, Once For 1 Occurrences Which drug do you need help dosing? prasugrel (EFFIENT) |
| () Maintenance Doses Only (Single Response) | |
| () clopidogrel (PLAVIX) 75 mg Maintenance Dose and aspirin EC 81 mg tablet - Start Tomorrow | |
| [] clopidogrel (PLAVIX) tablet | 75 mg, oral, daily, Starting S+1, Post-op |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| ticagrelor (BRILINTA) 90 mg Maintenance Dose and aspirin EC 81 mg tablet - Start 12 Hours from Now | |
| [] ticagrelor (BRILINTA) tablet | 90 mg, oral, 2 times daily, Starting H+12 Hours, Post-op |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| () prasugrel (EFFIENT) 10 mg Maintenance Dose and aspirin EC 81 mg tablet - Start Tomorrow | |
| [] prasugrel (EFFIENT) tablet + consult | "And" Linked Panel |
| [] prasugrel (EFFIENT) tablet | 10 mg, oral, daily, Starting S+1 |
| [] prasugrel (EFFIENT) consult | STAT, Once For 1 Occurrences Which drug do you need help dosing? prasugrel (EFFIENT) |
| [] aspirin (ECOTRIN) enteric coated tablet | 81 mg, oral, daily, Starting S+1, Post-op |
| () Anti-Platelet Contraindication | Routine, Until discontinued, Starting S Reason for "No" order: Post-op |
| | |

| () metoprolol tartrate (LOPRESSOR) tablet | 25 mg, oral, 2 times daily at 0600, 1800, Starting S+1, Post-op |
|---|---|
| | DO NOT administer if patient is on inotrope, vasopressor or |
| | has epicardial pacing |
| | HOLD parameters for this order: Hold Parameters requested |
| | HOLD for: 110 mmHg HOLD for Heart Rate LESS than: Other |
| | Please specify: 60 |
| | Contact Physician if: |
| () carvedilol (COREG) tablet | 3.125 mg, oral, 2 times daily at 0600, 1800, Starting S+1, |
| | Post-op |
| | DO NOT administer if heart rate is less than 60; systolic blood |
| | pressure is less than 110; on inotrope, vasopressor or has epicardial pacing |
| | HOLD parameters for this order: Hold Parameters requested |
| | HOLD for: 110 mmHg |
| | HOLD for: |
| | Contact Physician if: |
| Statin Therapy (Single Response) | |
| () simvastatin (ZOCOR) tablet | 40 mg, oral, nightly, Post-op |
| | Reduce to 20 mg daily if patient is on amiodarone. Do not give |
| | with grapefruit juice. |
| () atorvastatin (LIPITOR) tablet | 40 mg, oral, nightly, Post-op |
| | Do not give with grapefruit juice. |
| Respiratory Medications | |
| Scheduled | |
| [] Scheduled - albuterol (PROVENTIL) nebulizer solution | 2.5 mg, nebulization, Respiratory Therapy - every 6 hours, |
| | Post-op |
| (1.0.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1 | Aerosol Delivery Device: Hand-Held Nebulizer |
| [] Scheduled - ipratropium (ATROVENT) 0.02 % nebulizer solution | 0.5 mg, nebulization, Respiratory Therapy - every 6 hours, |
| Solution | Post-op Aerosol Delivery Device: Hand-Held Nebulizer |
|] PRN | riordoor Borrory Borros Franka Franka Franka |
| PRN - albuterol (PROVENTIL) nebulizer solution | 2.5 mg, nebulization, every 6 hours PRN, wheezing, Post-op Aerosol Delivery Device: Hand-Held Nebulizer |
| [] PRN - ipratropium (ATROVENT) 0.02 % nebulizer solution | 0.5 mg, nebulization, every 6 hours PRN, wheezing, Post-op Aerosol Delivery Device: Hand-Held Nebulizer |
| | |
| Multimodal Pain Management | |
| [X] pregabalin (LYRICA) capsule | 100 mg, oral, 2 times daily |
| [X] dexMEDEtomidine (PRECEDEX) infusion | 0.5 mcg/kg/hr, intravenous, continuous, Post-op |
| | Do not titrate without MD order; for postoperative pain. If |
| | needed for sedation, this order will need to be modified to the |
| | ICU sedation order to include titration parameters and dose range. Discontinue Dexmedetomidine (Precedex) IV infusion |
| | after extubation. Discontinue on postoperative day 1. |
| [X] acetaminophen (OFIRMEV) intravenous solution | 1,000 mg, intravenous, for 15 Minutes, every 6 hours, For 1 |
| | Doses, Post-op |
| | Maximum 3g/day. Total Tylenol/ acetaminophen dose (which |
| | includes, IV, PO or combination i.e. Norco, APAP etc) should |
| (V) contaminant on (TVI ENOL) tablet | not exceed 3g/day and 2g/day in case of cirrhotic patients. |
| [X] acetaminophen (TYLENOL) tablet | 1,000 mg, oral, every 6 hours PRN, mild pain (score 1-3), |
| A decian inophen (TTELIVOE) tablet | Starting Life Hours Doct on |
| A, acctaminophen (TTEENOE) tablet | Starting H+6 Hours, Post-op Total Tylenol/ acetaminophen dose (which includes TV PO or |
| A, acctaminophen (TTEENCE) tablet | Starting H+6 Hours, Post-op Total Tylenol/ acetaminophen dose (which includes, IV, PO or combination i.e. Norco, APAP etc) should not exceed 3g/day |

| [X] keTOROlac (TORadol) tablet | 30 mg, oral, every 6 hours, For 5 Days, Post-op Maximum 120mg/day in adults more than 50kg. Maximum 60mg/day in adults less than 50kg. |
|---|--|
| Moderate Break Through Pain (Single Response) | |
| (X) HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet | 1 tablet, oral, every 4 hours PRN, moderate pain (score 4-6), severe pain (score 7-10) Total Tylenol/ acetaminophen dose (which includes, IV, PO or combination i.e. Norco, APAP etc) should not exceed 3 g/d and 2 g/d in case of cirrhotic patients. |
| Severe Break Through Pain | |
| [X] HYDROcodone-acetaminophen (NORCO) 10-325 mg per tablet | 1 tablet, oral, every 4 hours PRN, severe pain (score 7-10), Post-op Total Tylenol/ acetaminophen dose (which includes, IV, PO or combination i.e. Norco, APAP etc) should not exceed 3 g/d and 2 g/d in case of cirrhotic patients. |
| [X] morPHINE injection | 2 mg, intravenous, every 1 hour prn, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed. |
| Antiemetics | |
| [] ondansetron (ZOFRAN) IV or Oral | "Or" Linked Panel |
| [] ondansetron ODT (ZOFRAN-ODT) disintegrating tablet | 4 mg, oral, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is able to tolerate oral medication. |
| [] ondansetron (ZOFRAN) 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 |
| promethazine (PHENERGAN) IV or Oral or Rectal | faster onset of action is required. "Or" Linked Panel |
| [] promethazine (PHENERGAN) 12.5 mg IV | 12.5 mg, intravenous, every 6 hours PRN, nausea, vomiting Post-op Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required. |
| [] promethazine (PHENERGAN) tablet | 12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-o Give if ondansetron (ZOFRAN) is ineffective and patient is able to tolerate oral medication. |
| [] promethazine (PHENERGAN) suppository | 12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral medication. |
| PUD Prophylaxis (Single Response) | |
| () famotidine (PEPCID) injection | 20 mg, intravenous, 2 times daily, Post-op |
| (X) pantoprazole (PROTONIX) EC tablet | 40 mg, oral, daily before breakfast, Post-op Indication(s) for Proton Pump Inhibitor (PPI) Therapy: |
| Bowel Care | |
| [X] Scheduled | |
| [X] Scheduled: polyethylene glycol (MIRALAX) packet - POD #1 | 17 g, oral, daily, Starting S+1, Post-op |
| [X] Docusate - Oral OR Nasogastric | "Or" Linked Panel |
| [X] docusate sodium (COLACE) capsule | 100 mg, oral, 2 times daily, Post-op Give if patient can tolerate oral medication |
| [X] docusate (COLACE) 50 mg/5 mL liquid | 100 mg, oral, 2 times daily, Post-op Give if patient has a nasogastric tube |
| [X] polyethylene glycol (MIRALAX) packet | 17 g, oral, daily, Post-op |

| [] As Needed: polyethylene glycol (MIRALAX) packet | 17 g, oral, daily PRN, constipation, Post-op RN may use second option based on the patient response to the first option attempted. |
|--|---|
| [] As Needed: Docusate - Oral OR Nasogastric | "Or" Linked Panel |
| [] docusate sodium (COLACE) capsule | 100 mg, oral, 2 times daily PRN, constipation, Post-op RN may use second option based on the patient response to the first option attempted. |
| [] docusate (COLACE) 50 mg/5 mL liquid | 100 mg, oral, 2 times daily PRN, constipation, Post-op RN may use second option based on the patient response to the first option attempted. Use if cannot swallow capsule. |
| [] sennosides-docusate sodium (SENOKOT-S) 8.6-50 mg per tablet | 1 tablet, oral, 2 times daily PRN, constipation, Post-op AS NEEDED AFTER FIRST BM |
| [] bisacodyl (DULCOLAX) suppository | 10 mg, rectal, daily PRN, constipation, Post-op FOR RECTAL USE ONLY. AS NEEDED TO MAINTAIN 3 BOWEL MOVEMENTS PER WEEK. DO NOT GIVE IF DIARRHEA NOTED. Administer if patient has not had a BM in 24 hours after oral therapy |
| emperature | |
| Acataminaphan aral par tuba ar ractal panal | "Or" Linked Banel |
| Acetaminophen oral, per tube or rectal panel Maximum of 3 grams of acetaminophen per day from all sour sources) | "Or" Linked Panel ces. (Cirrhosis patients maximum: 2 grams per day from all |
| Maximum of 3 grams of acetaminophen per day from all sour | |
| Maximum of 3 grams of acetaminophen per day from all sour sources) | ces. (Cirrhosis patients maximum: 2 grams per day from all 650 mg, oral, every 6 hours PRN, fever, GREATER than 100.4, Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day |

VTE

DVT Risk and Prophylaxis Tool (Single Response)

Low Risk Definition Moderate Risk Definition

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

Both pharmacologic AND mechanical prophylaxis must be addressed.

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

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

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

Central line Acute spinal cord injury with paresis

History of DVT or family history of VTE Multiple major traumas

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

Less than fully and independently ambulatory Acute ischemic stroke

Estrogen therapy History of PE

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

| () Low Risk of DVT | |
|--|---|
| [] Low Risk (Single Response) | |
| () Low risk of VTE | Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourgae early ambulation PACU & Post-op |
| () Moderate Risk of DVT - Surgical | |
| Address pharmacologic prophylaxis by selecting one of the follopharmacologic prophylaxis is contraindicated. | owing. Mechanical prophylaxis is optional unless |
| [] Moderate Risk | |
| [] Moderate risk of VTE | Routine, Once, PACU & Post-op |
| [] Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| () Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () enoxaparin (LOVENOX) injection (Single Response) | |
| () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| () enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min |

| () | 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): |
| () | 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 | 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 |
| | with high risk of bleeding, e.g. weight < 50kg and age > | AM, PACU & Post-op |
| | 75yrs) | Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| () | warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op |
| _ | | Indication: |
| () | Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| []_ | Mechanical Prophylaxis (Single Response) | |
| () | Contraindications exist for mechanical prophylaxis | Routine, Once |
| | | No mechanical VTE prophylaxis due to the following |
| | | contraindication(s): |
| | | PACU & Post-op |
| <u></u> | Place/Maintain cognential compression device | |
| () | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|) Mc | continuous oderate Risk of DVT - Non-Surgical | |
|) Mc | continuous | |
|) Mc Ad ph | continuous oderate Risk of DVT - Non-Surgical ddress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk | owing. Mechanical prophylaxis is optional unless |
|) Mo Ad pha | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE | |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | owing. Mechanical prophylaxis is optional unless Routine, Once, PACU & Post-op |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - | Routine, Once, PACU & Post-op Routine, Once |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | Routine, Once Routine, Once No pharmacologic VTE prophylaxis because: patient is |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follor armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follo armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follor armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follor armacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogarmacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response) | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogarmacologic prophylaxis is contraindicated. Moderate Risk Moderate risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response) | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogarmacologic prophylaxis is contraindicated. Moderate Risk Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response)) enoxaparin (LOVENOX) syringe) enoxaparin (LOVENOX) syringe - For Patients with CrCL | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting S 30 mg, subcutaneous, daily at 1700 (time critical), Starting |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogarmacologic prophylaxis is contraindicated. Moderate Risk Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response)) enoxaparin (LOVENOX) syringe) enoxaparin (LOVENOX) syringe - For Patients with CrCL | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting S 30 mg, subcutaneous, daily at 1700 (time critical), Starting S |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogarmacologic prophylaxis is contraindicated. Moderate Risk Moderate Risk of VTE Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response)) enoxaparin (LOVENOX) syringe) enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting S 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min |
|) Mc Ad ph: | continuous oderate Risk of DVT - Non-Surgical Idress pharmacologic prophylaxis by selecting one of the follogramacologic prophylaxis is contraindicated. Moderate Risk Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) Patient is currently receiving therapeutic anticoagulation Contraindications exist for pharmacologic prophylaxis enoxaparin (LOVENOX) injection (Single Response)) enoxaparin (LOVENOX) syringe) enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min) enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting S 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl |

| () 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 CrCI 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, 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, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|) High Risk of DVT - Surgical | |
| Address both pharmacologic and mechanical prophylaxis by or [] High Risk | rdering from Pharmacological and Mechanical Prophylaxis. |
| [] High risk of VTE | Routine, Once, PACU & Post-op |
| [] High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| () Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () an assertation (LOV/ENOV) in iterations (Circula Decompose) | 1 AOO & 1 OSt Op |
| () enoxaparin (LOVENOX) injection (Single Response) | |
| () enoxaparin (LOVENOX) injection (Single Response) () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| () enoxaparin (LOVENOX) syringe() enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min |
| () enoxaparin (LOVENOX) syringe() enoxaparin (LOVENOX) syringe - For Patients with CrCL | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |

| () 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. |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|) High Risk of DVT - Non-Surgical | |
| Address both pharmacologic and mechanical prophylaxis by or | dering from Pharmacological and Mechanical Prophylaxis. |
| | |
| [] High Risk | |
| [] High risk of VTE | Routine, Once, PACU & Post-op |
| [] High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | |
| () Patient is currently receiving therapeutic anticoagulation | |
| () Taken is carrently receiving the appeals anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. |
| | No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): |
| () Contraindications exist for pharmacologic prophylaxis | No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): |
| () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Response) | No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting |
| () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Response) () enoxaparin (LOVENOX) syringe () enoxaparin (LOVENOX) syringe - For Patients with CrCL | No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 40 mg, subcutaneous, daily at 1700 (time critical), Starting S 30 mg, subcutaneous, daily at 1700 (time critical), Starting S |

| () | 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): |
|---------------------|--|--|
| () | heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, PACU & Post-op |
| $\frac{\Box}{\Box}$ | heparin (porcine) injection (Recommended for patients | 5,000 Units, subcutaneous, every 12 hours, PACU & |
| () | with high risk of bleeding, e.g. weight < 50kg and age > | Post-op |
| | 75yrs) | Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| () | warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| () | Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] | Mechanical Prophylaxis (Single Response) | |
| () | Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| () Hig | gh Risk of DVT - Surgical (Hip/Knee) | |
| | dress both pharmacologic and mechanical prophylaxis by ord | dering from Pharmacological and Mechanical Prophylaxis. |
| [] | High Risk | |
| [] | High risk of VTE | Routine, Once, PACU & Post-op |
| | High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) | |
| () | Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () | Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () | apixaban (ELIQUIS) tablet | 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: |
| () | 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 |
| () | enoxaparin (LOVENOX) injection (Single Response) | 102 mg, orar, daily, clarting of 1, 1 7100 a 1 out op |
| () | | 40 mg, subcutaneous, daily at 0600 (time critical), Starting |
| () | | S+1 |
| () | | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 |
| () | enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min. |
| | enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. |
| () | enoxaparin (LOVENOX) syringe - For Patients weight 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 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |

| () 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 | > AM, PACU & Post-op |
| 75yrs) | Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs. |
| () rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (time critical), Starting S+1, PACU & Post-op |
| | To be Given on Post Op Day 1. Indications: |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
|] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |

DVT Risk and Prophylaxis Tool (Single Response)

Low Risk Definition Moderate Risk Definition

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

Both pharmacologic AND mechanical prophylaxis must be addressed.

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

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

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

Central line Acute spinal cord injury with paresis

History of DVT or family history of VTE Multiple major traumas

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

Less than fully and independently ambulatory Acute ischemic stroke

Estrogen therapy History of PE

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

| () Low Risk of DVT | |
|------------------------------------|--|
| [] Low Risk (Single Response) | |
| () Low risk of VTE | Routine, Once |
| | Low risk: Due to low risk, no VTE prophylaxis is needed. |
| | Will encourgae early ambulation |
| | PACU & Post-op |
| () Moderate Risk of DVT - Surgical | |

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

| Moderate Risk | |
|--|--|
| [] Moderate risk of VTE | Routine, Once, PACU & Post-op |
| [] Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| () Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () enoxaparin (LOVENOX) injection (Single Response) | |
| () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| () enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 For Patients with CrCL LESS than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| | 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): |
| () 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. |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|) Moderate Risk of DVT - Non-Surgical | |
| Address pharmacologic prophylaxis by selecting one of the follopharmacologic prophylaxis is contraindicated. | owing. Mechanical prophylaxis is optional unless |
| Moderate Risk | |
| [] Moderate risk of VTE | Routine, Once, PACU & Post-op |
| [] Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | |

| () Patient is currently receiving therapeutic anticoagulation | |
|--|--|
| () Tallett is currently receiving therapeduc anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () enoxaparin (LOVENOX) injection (Single Response) | |
| () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| () enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 1700 (time critical), Starting S For Patients with CrCL LESS than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight | 30 mg, subcutaneous, 2 times daily, Starting S |
| between 100-139 kg and CrCl GREATER than 30 mL/min | For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| () 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): |
| () heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, 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, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. |
| | weight LESS than 50kg and age GREATER than 75yrs. |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
|) High Risk of DVT - Surgical | |
| Address both pharmacologic and mechanical prophylaxis by ord | dering from Pharmacological and Mechanical Prophylaxis. |
| [] High Risk | |
| [] High risk of VTE | Routine, Once, PACU & Post-op |
| [] High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) | |
| () Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| | Routine, Once |

| () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
|---|---|
| () enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min | 30 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 |
| | For Patients with CrCL LESS than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl |
| | GREATER than 30 mL/min |
| () enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| () 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. |
| () warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |
| High Risk of DVT - Non-Surgical | |
| Address both pharmacologic and mechanical prophylaxis by ord | dering from Pharmacological and Mechanical Prophylaxis. |
|] High Risk [] High risk of VTE | Routine, Once, PACU & Post-op |
| High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) | Routine, Once, i Aco & i ost-op |
| () Patient is currently receiving therapeutic anticoagulation | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op |
| () Contraindications exist for pharmacologic prophylaxis | Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () enoxaparin (LOVENOX) injection (Single Response) | |
| () enoxaparin (LOVENOX) syringe | 40 mg, subcutaneous, daily at 1700 (time critical), Starting S |
| () enoxaparin (LOVENOX) syringe - For Patients with CrCL | 30 mg, subcutaneous, daily at 1700 (time critical), Starting |

| enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min |
|--|--|
| () enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min | 40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| () 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): |
| () heparin (porcine) injection | 5,000 Units, subcutaneous, every 8 hours, 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, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. |
| () warfarin (COUMADIN) tablet | weight LESS than 50kg and age GREATER than 75yrs. oral, daily at 1700 (time critical), PACU & Post-op Indication: |
| () Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
| [] Mechanical Prophylaxis (Single Response) | |
| () Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| () Place/Maintain sequential compression device | Routine, Continuous, PACU & Post-op |
| continuous | |
| | |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore | dering from Pharmacological and Mechanical Prophylaxis. |
| High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore High Risk | |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore [] High Risk [] High risk of VTE | dering from Pharmacological and Mechanical Prophylaxis. Routine, Once, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore [] High Risk | Routine, Once, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore [] High Risk | |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore [] High Risk | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ore [] High Risk | Routine, Once, PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) () Patient is currently receiving therapeutic anticoagulation () Contraindications exist for pharmacologic prophylaxis () apixaban (ELIQUIS) tablet | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) () Patient is currently receiving therapeutic anticoagulation () Contraindications exist for pharmacologic prophylaxis () apixaban (ELIQUIS) tablet () aspirin chewable tablet | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) () Patient is currently receiving therapeutic anticoagulation () Contraindications exist for pharmacologic prophylaxis () apixaban (ELIQUIS) tablet () aspirin chewable tablet () aspirin (ECOTRIN) enteric coated tablet | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) () Patient is currently receiving therapeutic anticoagulation () Contraindications exist for pharmacologic prophylaxis () apixaban (ELIQUIS) tablet () aspirin chewable tablet () aspirin (ECOTRIN) enteric coated tablet () enoxaparin (LOVENOX) injection (Single Response) | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: 162 mg, oral, daily, Starting S+1, PACU & Post-op 162 mg, oral, daily, Starting S+1, PACU & Post-op |
| () High Risk of DVT - Surgical (Hip/Knee) Address both pharmacologic and mechanical prophylaxis by ord [] High Risk [] High risk of VTE [] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) () Patient is currently receiving therapeutic anticoagulation () Contraindications exist for pharmacologic prophylaxis () apixaban (ELIQUIS) tablet () aspirin chewable tablet () aspirin (ECOTRIN) enteric coated tablet () enoxaparin (LOVENOX) injection (Single Response) () enoxaparin (LOVENOX) syringe | Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: PACU & Post-op Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op 2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: 162 mg, oral, daily, Starting S+1, PACU & Post-op 162 mg, oral, daily, Starting S+1, PACU & Post-op 40 mg, subcutaneous, daily at 0600 (time critical), Starting S+1 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time |

| () | enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min | 30 mg, subcutaneous, 2 times daily at 0600, 1800 (time critical), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. |
|-----|---|--|
| () | enoxaparin (LOVENOX) syringe - For Patients weight 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 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min |
| () | 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. |
| | rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission | 10 mg, oral, daily at 0600 (time critical), Starting S+1, PACU & Post-op To be Given on Post Op Day 1. Indications: |
| () | warfarin (COUMADIN) tablet | oral, daily at 1700 (time critical), Starting S+1, PACU & Post-op Indication: |
| () | Pharmacy consult to manage warfarin (COUMADIN) | STAT, Until discontinued, Starting S Indication: |
|] N | Mechanical Prophylaxis (Single Response) | |
| | Contraindications exist for mechanical prophylaxis | Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op |
| | Place/Maintain sequential compression device continuous | Routine, Continuous, PACU & Post-op |

| Labs | |
|------------|--|
| Labs Today | |

| Labs Today | |
|--|------------------------------------|
| [] Lactic acid level | Once, Post-op |
| [X] Basic metabolic panel | Once, Post-op |
| [X] CBC with platelet and differential | Once, Post-op |
| [] Magnesium level | Once, Post-op |
| [] Phosphorus level | Once, Post-op |
| [] Calcium level | Once, Post-op |
| [] Ionized calcium | Once, Post-op |
| [] Prothrombin time with INR | Once, Post-op |
| [] Partial thromboplastin time | Once, Post-op |
| [] Platelet function P2Y12 | Once, Post-op |
| [] Platelet mapping | Once |
| | Anticoagulant Therapy: |
| | Diagnosis: |
| | Fax Number (For TEG Graph Result): |
| | Post-op |
| [] Troponin | Once, Post-op |
| [] B natriuretic peptide | Once, Post-op |
| [] Anti Xa, unfractionated | Once, Post-op |
| [] Fibrinogen | Once, Post-op |

| [] Cortisol level, random | Once, Post-op |
|--|---|
| [] Type and screen | Once, Post-op |
| [] Blood gas, arterial | Once, Post-op |
| Labs Every 8 hours x 3 | |
| [] Troponin | Now then every 8 hours For 3 Occurrences, Post-op |
| DIC Panel | |
| [] Partial thromboplastin time | Once, Post-op |
| Prothrombin time with INR | Once, Post-op |
| [] Fibrinogen | Once, Post-op |
| [] D-dimer | Once, Post-op |
| Labs Every AM x 3 Days | |
| [] CBC hemogram | AM draw repeats For 3 Occurrences, Post-op |
| Basic metabolic panel | AM draw repeats For 3 Occurrences, Post-op |
| Magnesium level | AM draw repeats For 3 Occurrences, Post-op |
| [] Phosphorus level | AM draw repeats For 3 Occurrences, Post-op |
| [] Ionized calcium | AM draw repeats For 3 Occurrences, Post-op |
| Cardiology | |
| Cardiology | |
| [X] ECG 12 lead - Once | Routine, Once |
| • | Clinical Indications: Post-Op Surgery |
| | Interpreting Physician: |
| | Post operative, Post-op |
| [] ECG 12 lead - Daily starting tomorrow | Routine, Daily, Starting S+1 For 3 Occurrences |
| | Clinical Indications: Post-Op Surgery |
| | Interpreting Physician: |
| 1. E. L | Post-op |
| Echocardiogram complete w contrast and 3D if needed | Routine, 1 time imaging, Post-op |
| | |
| Imaging | |
| | |
| X-Ray [X] Chest 1 Vw Portable | Routine, 1 time imaging For 1 Occurrences, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) | Routine, Daily imaging For 3 Occurrences, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) | Routine, Daily imaging For 3 Occurrences, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op |
| [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory Respiratory | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory Respiratory [X] Encourage deep breathing and coughing | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory Respiratory [X] Encourage deep breathing and coughing | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable (after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry [] Positive Expiratory (PEP) Device | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op |
| X-Ray [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op Routine, Continuous |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable(after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry [] Positive Expiratory (PEP) Device | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op Routine, Continuous Device 1: |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable (after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry [] Positive Expiratory (PEP) Device | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op Routine, Continuous Device 1: Device 2: |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable (after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry [] Positive Expiratory (PEP) Device | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op Routine, Continuous Device 1: Device 2: Device 3: |
| [X] Chest 1 Vw Portable [] Chest 1 Vw Portable (Daily) [X] Chest 1 Vw Portable (after chest tube removal) Ultrasound [] PV carotid duplex bilateral Respiratory [X] Encourage deep breathing and coughing [X] Incentive spirometry [] Positive Expiratory (PEP) Device | Routine, Daily imaging For 3 Occurrences, Post-op Routine, Conditional Frequency For 1 Occurrences After chest tube removal, Post-op Routine, 1 time imaging, Post-op Routine, Every hour, Post-op Routine, Once 10 x every hour while awake, Post-op Routine, Once Twenty (20) times every hour while awake, Post-op Routine, Continuous Device 1: Device 2: |

| Consults | |
|--|--|
| For Physician Consult orders use sidebar | |
| · | |
| Ancillary Consults | |
| | O !(D D' ! D! ' |
| [] Consult to Case Management | Consult Reason: Discharge Planning |
| [1] Consult to Conicl World | Post-op |
| [] Consult to Social Work | Reason for Consult: |
| I. Consult DT and treat | Post-op |
| [] Consult PT eval and treat | Special Instructions: |
| [1] Consult DT waved asse | Weight Bearing Status: |
| [] Consult PT wound care | Special Instructions: Location of Wound? |
| | |
| [] Consult OT eval and treat | Post-op |
| [] Consult OT eval and treat | Special Instructions: Weight Bearing Status: |
| [] Consult to Nutrition Services | Reason For Consult? |
| [] Consult to Nutrition Services | Purpose/Topic: |
| | Post-op |
| [] Consult to Spiritual Care | Reason for consult? |
| [] Consult to Spiritual Care | Post-op |
| Consult to Speech Language Pathology | Routine, Once |
| [] Consult to Speech Language Fathology | Reason for consult: |
| | Post-op |
| [] Consult to Wound Ostomy Care nurse | Reason for consult: |
| [1] Contain to Tround Colomy Care names | Reason for consult: |
| | Reason for consult: |
| | Reason for consult: |
| | Consult for NPWT: |
| | Reason for consult: |
| | Post-op |
| [] Consult to Respiratory Therapy | Reason for Consult? |
| | Post-op |

Additional Orders