

General

Common Present on Admission Diagnosis

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

Elective Outpatient, Observation, or Admission (Single Response)

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

Other

Admission or Observation (Single Response)

Patient has active outpatient status order on file

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

Other

Admission (Single Response)

Patient has active status order on file

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

Transfer (Single Response)

Patient has active inpatient status order on file

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

Isolation

- | | |
|---|-----------------------|
| <input type="checkbox"/> Airborne isolation status | |
| <input type="checkbox"/> Airborne isolation status | Details |
| <input type="checkbox"/> Mycobacterium tuberculosis by PCR - If you suspect Tuberculosis, please order this test for rapid diagnostics. | Once, Sputum, Post-op |
| <input type="checkbox"/> Contact isolation status | Details |
| <input type="checkbox"/> Droplet isolation status | Details |
| <input type="checkbox"/> Enteric isolation status | Details |
| <input type="checkbox"/> Other | |

Precautions

- | | |
|---|--|
| <input type="checkbox"/> Aspiration precautions | Post-op |
| <input type="checkbox"/> Fall precautions | Increased observation level needed:
if (answer = Yes)
Level:
For:
Time:
Post-op |

<input type="checkbox"/>	Latex precautions	Post-op
<input type="checkbox"/>	Seizure precautions	Increased observation level needed: if (answer = Yes) Level: For: Time: Post-op
<input type="checkbox"/>	Other	

Nursing

Vitals

<input type="checkbox"/>	Vital signs - T/P/R/BP	Routine, Per unit protocol, Starting S On arrival and then routine, Post-op
<input type="checkbox"/>	Other	

Activity

<input type="checkbox"/>	Head of bed	Routine, Until discontinued, Starting S Head of bed: 30 degrees if (answer = other degrees (specify)) Specify: Post-op
<input type="checkbox"/>	Strict bed rest	Routine, Until discontinued, Starting S For 24 Hours For 24 hours PostOp, Post-op
<input type="checkbox"/>	Up in chair	Routine, Until discontinued, Starting S+1 Specify: Up in chair if (answer = Up in chair) Additional modifier: if (answer = Other activity (specify)) Other: Additional modifier: Starting 24 hours post-operative., Post-op
<input type="checkbox"/>	Other	

Nursing

<input type="checkbox"/>	Insert feeding tube	Routine, Once For 1 Occurrences Insert Dobhoff tube, Post-op
<input type="checkbox"/>	Strict intake and output	Routine, Every 8 hours Per floor protocol, Post-op
<input type="checkbox"/>	Foley catheter care	Routine, Until discontinued, Starting S Orders: Maintain, to gravity Post-op
<input type="checkbox"/>	Nasogastric tube maintenance	Routine, Until discontinued, Starting S Tube Care Orders: To Low Intermittent Suction Remove after extubation, Post-op

<input type="checkbox"/> Orogastric tube maintenance	Routine, Until discontinued, Starting S Tube Care Orders: To Low Intermittent Suction Remove after extubation, Post-op
<input type="checkbox"/> Apply warming blanket	Routine, As needed As needed to raise body temperature to 98.6 Fahrenheit, Post-op
<input type="checkbox"/> Reinforce dressing	Routine, As needed Reinforce with: Reinforce dressing as needed., Post-op
<input type="checkbox"/> Hemodynamic Monitoring	Routine, Every hour Measure: Other if (answer = Other) Other: Other: Swan Ganz to monitor, Recalibrate SV02 every morning. Record SV02 every 1 hour. DO NOT WEDGE SWAN. Post-op
<input type="checkbox"/> Chest tube to continuous suction	Routine, Until discontinued, Starting S Level of suction: 20 cm H2O Chest tube to 20 centimeter water pressure., Post-op
<input type="checkbox"/> Neurological assessment	Routine, Daily Assessment to Perform: if (answer = Spinal exams) Perform: Area: Post-op
<input type="checkbox"/> Hold sedation every morning coordinated with CV intensivist to assess neurological status	Routine, Until discontinued, Starting S, Post-op
<input type="checkbox"/> All blood products must be irradiated and leukocyte reduced	Routine, Until discontinued, Starting S, Post-op
<input type="checkbox"/> Blood products must be CMV negative if donor and recipients are CMV negative	Routine, Until discontinued, Starting S, Post-op
<input type="checkbox"/> Other	

Notify Physician

<input type="checkbox"/> Notify Physician for vitals:	Routine, Until discontinued, Starting S Temperature greater than: 100.1 Temperature less than: Systolic BP greater than: 160 Systolic BP less than: 90 Diastolic BP greater than: 110 Diastolic BP less than: 40 MAP less than: 60 Heart rate greater than (BPM): 120 Heart rate less than (BPM): 50 Respiratory rate greater than: 25 Respiratory rate less than: 8 SpO2 less than: 88
<input type="checkbox"/> Notify CV Intensivist for critical labs, vomiting, GI bleed, cardiac arrhythmias, chest pain	Routine, Until discontinued, Starting S, Post-op

<input type="checkbox"/>	CV Intensivist if chest tube drainage greater than 100 millimeters in 2 hours.	Routine, Until discontinued, Starting S, Post-op
<input type="checkbox"/>	Other	

Diet

<input type="checkbox"/>	NPO	Diet effective now, Starting S NPO: Pre-Operative fasting options: if (answer = Other) Specify: Post-op
<input type="checkbox"/>	Other	

IV Fluids

IV Fluids (Single Response)

<input type="checkbox"/>	sodium chloride 0.9 % bolus	500 mL, intravenous, for 30 Minutes, once, For 1 Doses, Post-op
<input type="checkbox"/>	sodium chloride 0.9 % infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	sodium chloride 0.45 % infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	dextrose 5%-0.45% sodium chloride infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	dextrose 5%-0.9% sodium chloride infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	sodium chloride 0.9 % with potassium chloride 20 mEq/L infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	dextrose 5 % and sodium chloride 0.45 % with potassium chloride 20 mEq/L infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	dextrose 5 % and sodium chloride 0.9 % with potassium chloride 20 mEq/L infusion	75 mL/hr, intravenous, at 100 mL/hr, continuous, Post-op
<input type="checkbox"/>	sodium chloride 0.45 % 1,000 mL with sodium bicarbonate 75 mEq/L infusion	75 mL/hr, intravenous, continuous, Post-op
<input type="checkbox"/>	Other	

Medications

Pharmacy Consult

<input type="checkbox"/>	Pharmacy consult to manage dose adjustments for renal function	STAT, Until discontinued, Starting S Adjust dose for:
<input type="checkbox"/>	Other	

Restricted Medications

<input type="checkbox"/>	No ketorolac (Toradol)	STAT, Until discontinued, Starting S Reason for "No" order: Status Post Lung Transplant Post-op
<input type="checkbox"/>	No NSAIDs EXcluding aspirin	STAT, Until discontinued, Starting S Reason for "No" order: Post-op
<input type="checkbox"/>	Other	

Induction Therapy

<input type="checkbox"/> basiliximab (SIMULECT) IVPB - POD #4	20 mg, intravenous, for 30 Minutes, once, S+4 at 11:00 AM, For 1 Doses, Post-op Administer on POD #4
<input type="checkbox"/> Other	

Immunosuppressants

<input type="checkbox"/> Immunosuppression Therapy: Option 1 - methylPREDNISolone and predniSONE	"Followed by" Linked Panel
<input type="checkbox"/> methylPREDNISolone (Solu-MEDROL) IV Push - POD #1 and 2	2 mg/kg, intravenous, daily, Starting S+1, For 2 Doses, Post-op
<input type="checkbox"/> methylPREDNISolone (Solu-MEDROL) IV Push - POD #3 and 4	1.5 mg/kg, intravenous, daily, Starting S+3, For 2 Doses, Post-op
<input type="checkbox"/> methylPREDNISolone (Solu-MEDROL) IV Push - POD #5 and 6	1 mg/kg, intravenous, daily, Starting S+5, For 2 Doses, Post-op
<input type="checkbox"/> predniSONE (DELTASONE) tablet - POD #7 and 8	40 mg, oral, daily, Starting S+7, For 2 Doses, Post-op
<input type="checkbox"/> predniSONE (DELTASONE) tablet - POD #9 and 10	20 mg, oral, daily, Starting S+9, For 2 Doses, Post-op
<input type="checkbox"/> predniSONE (DELTASONE) tablet - POD #11	10 mg, oral, daily, Starting S+11, Post-op
<input type="checkbox"/> Immunosuppression Therapy: Option 2 - mycophenolate IVPB	
<input type="checkbox"/> Immunosuppression Therapy: Option 3 - mycophenolate (CELLCEPT) IVPB	1,000 mg, intravenous, for 2 Hours, 2 times daily, Starting S, Post-op
<input type="checkbox"/> Immunosuppression Therapy: Option 3 - tacrolimus NG Tube or Oral and cyclosporine NG Tube or Oral (Single Response)	
<input type="checkbox"/> tacrolimus (PROGRAF) 0.5 mg/ml oral suspension - POD #1	Nasogastric, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1, Post-op Clamp Nasogastric tube times 1 hour.
<input type="checkbox"/> tacrolimus (PROGRAF) capsule - POD #1	sublingual, 2 times daily at 0600, 1800, Starting S+1, Post-op Open the capsule and put the contents under the tongue.
<input type="checkbox"/> cycloSPORINE (NEORAL) solution - POD #1	Nasogastric, 2 times daily at 0600, 1800, Starting S+1, Post-op Clamp Nasogastric tube times 1 hour.
<input type="checkbox"/> cycloSPORINE (NEORAL) capsule - POD #1	oral, 2 times daily at 0600, 1800, Starting S+1, Post-op
<input type="checkbox"/> Other	

Pneumocystis Prophylaxis

<input type="checkbox"/> sulfamethoxazole-trimethoprim (BACTRIM DS) Options (Single Response)	
<input type="checkbox"/> sulfamethoxazole-trimethoprim (BACTRIM DS) 800-160 mg tablet	1 tablet, oral, user specified, Starting S+5, Post-op Give on PostOp Day 5. Type of Therapy: New Anti-Infective Order if (answer = New Anti-Infective Order) Reason for Therapy: if (answer = Other) Specify: if (answer = Bacterial Infection Suspected) Indication: if (answer = Other) Specify: if (answer = Bacterial Infection Documented) Indication: if (answer = Other)

Specify:
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

sulfamethoxazole-trimethoprim (BACTRIM) 200-40 mg/5 mL suspension 20 mL, Nasogastric, user specified, Starting S+5, Post-op
Give on PostOp Day 5.
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)
Reason for Therapy:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

atovaquone (MEPRON) suspension - If Known or Suspected Sulfa Allergy - POD #5 750 mg, Nasogastric, 2 times daily, Starting S+5
If Known or Suspected Sulfa Allergy

Other

Antivirals (Single Response)

valGANCiclovir (VALCYTE) 50 mg/mL oral solution 450 mg, oral, 2 times daily, Post-op
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)
Reason for Therapy:
if (answer = Other)

Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:
Reason for Therapy: Medical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

() ganciclovir (CYTOVENE) IVPB - POD #0

5 mg/kg, intravenous, daily, Starting S, Post-op
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)
Reason for Therapy:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:
Reason for Therapy: Medical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

() acyclovir (ZOVIRAX) IV

5 mg/kg, intravenous, every 8 hours, Post-op
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)

Reason for Therapy:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

Reason for Therapy: Medical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

() acyclovir (ZOVIRAX)

200 mg, oral, 2 times daily, Post-op
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)
Reason for Therapy:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:
Reason for Therapy: Medical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

[] Other

Antifungal

[] nystatin (MYCOSTATIN) 100,000 unit/mL suspension

5 mL, oral, 4 times daily, Post-op
Paint mouth with swab while intubated. Once extubated, convert to swish and swallow.

Type of Therapy: New Anti-Infective Order

if (answer = New Anti-Infective Order)

Reason for Therapy:

if (answer = Other)

Specify:

if (answer = Bacterial Infection Suspected)

Indication:

if (answer = Other)

Specify:

if (answer = Bacterial Infection Documented)

Indication:

if (answer = Other)

Specify:

Reason for Therapy: Surgical Prophylaxis

if (answer = Other)

Specify:

if (answer = Bacterial Infection Suspected)

Indication:

if (answer = Other)

Specify:

if (answer = Bacterial Infection Documented)

Indication:

if (answer = Other)

Specify:

[] voriconazole (VFEND) tablet - POD #1

200 mg, oral, 2 times daily, Starting S+1, Post-op
Crush tablet to make suspension if patient is unable to swallow.

Type of Therapy: New Anti-Infective Order

if (answer = New Anti-Infective Order)

Reason for Therapy:

if (answer = Other)

Specify:

if (answer = Bacterial Infection Suspected)

Indication:

if (answer = Other)

Specify:

if (answer = Bacterial Infection Documented)

Indication:

if (answer = Other)

Specify:

Reason for Therapy: Surgical Prophylaxis

if (answer = Other)

Specify:

if (answer = Bacterial Infection Suspected)

Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

[] micafungin (MYCAMINE) IVPB

100 mg, intravenous, for 1 Hours, every 24 hours, For 2 Doses, Post-op
RESTRICTED to Infectious Diseases (ID), Solid Organ Transplant (SOT), Bone
Marrow Transplant (BMT), and Hematology/Oncology (Heme/Onc) specialists.
Are you an ID, SOT, BMT, or Heme/Onc specialist or ordering on behalf of one?
YES, I am an approved provider
if (answer = I am ordering on behalf of an approved provider)
Name of Approved Provider:
if (answer = NO)
HM Policy Alert:
if (answer = Formulary policy override (pharmacist use only))
Provide name of secondary pharmacist who provided authorization and open
a "Formulary Policy Override" i-Vent:
Type of Therapy: New Anti-Infective Order
if (answer = New Anti-Infective Order)
Reason for Therapy:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:

[] ipratropium (ATROVENT) 0.02 % nebulizer solution FOLLOWED BY
amphotericin B liposome (AMBISOME) 50 mg inhalation suspension

"Followed by" Linked Panel

<input type="checkbox"/> ipratropium (ATROVENT) 0.02 % nebulizer solution	0.5 mg, nebulization, Respiratory Therapy - every 4 hours, For 7 Days, Post-op Aerosol Delivery Device: if (answer = Intrapulmonary Percussive Ventilation (Meta-Neb Device)) Meta-Neb Indications:
<input type="checkbox"/> amphotericin B liposome (AMBISOME) 50 mg in water for injection, sterile (PF) 6.25 mL inhalation suspension	50 mg, inhalation, Respiratory Therapy - Daily, Post-op RESTRICTED to Infectious Diseases (ID), Solid Organ Transplant (SOT), Bone Marrow Transplant (BMT), and Hematology/Oncology (Heme/Onc) specialists. Are you an ID, SOT, BMT, or Heme/Onc specialist or ordering on behalf of one? if (answer = I am ordering on behalf of an approved provider) Name of Approved Provider: if (answer = NO) HM Policy Alert: if (answer = Formulary policy override (pharmacist use only)) Provide name of secondary pharmacist who provided authorization and open a "Formulary Policy Override" i-Vent: [amphotericin B liposome]Reason for Therapy: if (answer = Other) Specify: if (answer = Fungal Infection Documented) Indication: if (answer = Other) Specify: Authorizing ID: if (answer = Other) Specify: if (answer = Fungal Infection Suspected) Indication: if (answer = Other) Specify: Authorizing ID: if (answer = Other) Specify:
<input type="checkbox"/> Other	

PostOperative Antibiotics - Gram Positive coverage (Single Response)

Gram Positive Coverage Antibiotics (Single Response)

- | | |
|---|--|
| <p>() vancomycin (VANCOGIN) IV - Administer 1 hour PRIOR to skin incision.</p> | <p>15 mg/kg, intravenous, every 12 hours, For 7 Days, Post-op
Administer 1 hour PRIOR to skin incision.
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:</p> |
| <p>() linezolid in dextrose 5% (ZYVOX) IVPB - For Known/Suspected Allergies or Suspected Drug-Resistant Organism to Vancomycin</p> | <p>600 mg, intravenous, for 60 Minutes, every 12 hours, For 7 Days, Post-op
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:</p> |

[] Other

PostOperative Antibiotics - Gram Negative coverage (Single Response)

() Gram Negative Coverage Antibiotics (Single Response)

Select ONE of the following:

- | | |
|---|--|
| <p>() piperacillin-tazobactam (ZOSYN) IV</p> | <p>3.375 g, intravenous, every 6 hours, For 7 Days, Post-op
Reason for Therapy: Surgical Prophylaxis
if (answer = Other)
Specify:
if (answer = Bacterial Infection Suspected)
Indication:
if (answer = Other)
Specify:
if (answer = Bacterial Infection Documented)
Indication:
if (answer = Other)
Specify:</p> |
|---|--|

<input type="checkbox"/> ceFEPime (MAXIPIME) IV	2 g, intravenous, every 8 hours, For 7 Days, Post-op Reason for Therapy: Surgical Prophylaxis if (answer = Other) Specify: if (answer = Bacterial Infection Suspected) Indication: if (answer = Other) Specify: if (answer = Bacterial Infection Documented) Indication: if (answer = Other) Specify:
<input type="checkbox"/> meropenem (MERREM) IV	500 mg, intravenous, every 6 hours, For 7 Days, Post-op Reason for Therapy: Surgical Prophylaxis if (answer = Other) Specify: if (answer = Bacterial Infection Suspected) Indication: if (answer = Other) Specify: if (answer = Bacterial Infection Documented) Indication: if (answer = Other) Specify:
<input type="checkbox"/> aztreonam (AZACTAM) IV	2 g, intravenous, every 8 hours, For 7 Days, Post-op For Known or Suspected Penicillin Allergy Reason for Therapy: Surgical Prophylaxis if (answer = Other) Specify: if (answer = Bacterial Infection Suspected) Indication: if (answer = Other) Specify: if (answer = Bacterial Infection Documented) Indication: if (answer = Other) Specify:
<input type="checkbox"/> Other	

Stress Ulcer Prophylaxis (Single Response)

<input type="checkbox"/> pantoprazole (PROTONIX) injection	40 mg, intravenous, daily at 0600, Post-op Indication(s) for Proton Pump Inhibitor (PPI) Therapy: if (answer = Other (Specify)) Specify:
<input type="checkbox"/> famotidine (PEPCID) injection	20 mg, intravenous, 2 times daily, Post-op
<input type="checkbox"/> Other	

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

(adjust dose for renal/liver function and age)

<input type="checkbox"/> acetaminophen (TYLENOL) tablet OR oral solution	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> acetaminophen (TYLENOL) tablet	650 mg, oral, every 6 hours PRN, mild pain (score 1-3), Post-op Maximum of 3 grams of acetaminophen per day from all sources. Give the tablet if the patient can tolerate oral medication. (Cirrhosis patients maximum: 2 grams per day from all sources)
<input type="checkbox"/> acetaminophen (TYLENOL)suspension	650 mg, oral, every 6 hours PRN, mild pain (score 1-3), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Use if patient cannot tolerate oral tablet.
<input type="checkbox"/> Other	

PRN Oral for Moderate Pain (Pain Score 4-6): For Patients LESS than 65 years old (Single Response)

(adjust dose for renal/liver function and age)

<input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) 300-30 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if patient is able to tolerate oral medication.
<input type="checkbox"/> acetaminophen-codeine 300 mg-30 mg /12.5 mL solution	12.5 mL, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient cannot swallow tablet.
<input type="checkbox"/> HYDROcodone-acetaminophen 5/325 (NORCO) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6)
<input type="checkbox"/> HYDROcodone-acetaminophen (HYCET) 2.5-108.3 mg/5 mL solution	10 mL, oral, every 6 hours PRN, moderate pain (score 4-6)
<input type="checkbox"/> HYDROcodone-acetaminophen 7.5/325 (NORCO) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 7.5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if patient is able to tolerate oral medication.
<input type="checkbox"/> HYDROcodone-acetaminophen (HYCET) 7.5-325 mg/15 mL solution	15 mL, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient cannot swallow tablet.

<input type="checkbox"/> HYDROcodone-acetaminophen 10/325 (NORCO) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO 10-325) 10-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if patient is able to tolerate oral medication.
<input type="checkbox"/> HYDROcodone-acetaminophen (HYCET) 7.5-325 mg/15 mL solution	20 mL, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient can not swallow tablet.
<input type="checkbox"/> traMADol (ULTRAM) tablet - For eGFR LESS than 30 mL/min, change frequency to every 12 hours)	50 mg, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op (Max Daily dose not to exceed 200 mg/day).
Give if patient is able to tolerate oral medication	
<input type="checkbox"/> Other	

PRN Oral for Moderate Pain (Pain Score 4-6): For Patients GREATER than 65 years old (Single Response)
(adjust dose for renal/liver function and age)

<input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> acetaminophen-codeine (TYLENOL #3) 300-30 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if patient is able to tolerate oral medication.
<input type="checkbox"/> acetaminophen-codeine 300 mg-30 mg /12.5 mL solution	12.5 mL, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient cannot swallow tablet.
<input type="checkbox"/> HYDROcodone-acetaminophen 5/325 (NORCO) tablet OR elixir	"Or" Linked Panel
Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)	
<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6)
<input type="checkbox"/> HYDROcodone-acetaminophen (HYCET) 2.5-108.3 mg/5 mL solution	10 mL, oral, every 6 hours PRN, moderate pain (score 4-6)
<input type="checkbox"/> traMADol (ULTRAM) tablet - For eGFR LESS than 30 mL/min, change frequency to every 12 hours)	25 mg, oral, every 6 hours PRN, moderate pain (score 4-6), Post-op (Max Daily dose not to exceed 200 mg/day). Give if patient can tolerate oral medication.
<input type="checkbox"/> Other	

PRN IV for Moderate Pain (Pain Score 4-6): For Patients LESS than 65 years old (Single Response)
If you select a PCA option you will not be allowed to also order IV PRN pain medications from this section.
(adjust dose for renal/liver function and age)

<input type="checkbox"/> fentaNYL (SUBLIMAZE) injection	25 mcg, intravenous, every 2 hour PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> morphine 2 mg/mL injection	2 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> HYDROmorphine (DILAUDID) injection	0.5 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> Other	

PRN IV for Moderate Pain (Pain Score 4-6): For Patients GREATER than 65 years old (Single Response)

If you select a PCA option you will not be allowed to also order IV PRN pain medications from this section.
(adjust dose for renal/liver function and age)

<input type="checkbox"/> fentaNYL (SUBLIMAZE) injection	12.5 mcg, intravenous, every 2 hour PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> morphine 2 mg/mL injection	1 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> HYDROmorphine (DILAUDID) injection	0.2 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> Other	

PRN Oral for Severe Pain (Pain Score 7-10): For Patients LESS than 65 years old (Single Response)

(adjust dose for renal/liver function and age)

<input type="checkbox"/> HYDROmorphine (DILAUDID) tablet	2 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> morphine (MSIR) tablet	15 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> oxyCODONE (ROXICODONE) immediate release tablet	10 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> Other	

PRN Oral for Severe Pain (Pain Score 7-10): For Patients GREATER than 65 years old (Single Response)

(adjust dose for renal/liver function and age)

<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO) 7.5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> HYDROcodone-acetaminophen (NORCO 10-325) 10-325 mg per tablet	1 tablet, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> HYDROmorphine (DILAUDID) tablet	2 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> morphine (MSIR) tablet	15 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> oxyCODONE (ROXICODONE) immediate release tablet	5 mg, oral, every 6 hours PRN, severe pain (score 7-10), Post-op Give if patient is able to tolerate oral medication
<input type="checkbox"/> Other	

PRN IV for Severe Pain (Pain Score 7-10): For Patients LESS than 65 years old (Single Response)

If you select a PCA option you will not be allowed to also order IV PRN pain medications from this section.
(adjust dose for renal/liver function and age)

<input type="checkbox"/> fentaNYL (SUBLIMAZE) injection	50 mcg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> morphine injection	4 mg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> HYDROmorphine (DILAUDID) injection	0.8 mg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> Other	

PRN IV for Severe Pain (Pain Score 7-10): For Patients GREATER than 65 years old (Single Response)

If you select a PCA option you will not be allowed to also order IV PRN pain medications from this section.
(adjust dose for renal/liver function and age)

<input type="checkbox"/> fentaNYL (SUBLIMAZE) injection	25 mcg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> morphine injection	2 mg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> HYDROmorphine (DILAUDID) injection	0.5 mg, intravenous, every 3 hours PRN, severe pain (score 7-10), Post-op Use if patient is unable to swallow or faster onset is needed
<input type="checkbox"/> Other	

Bowel Care

<input type="checkbox"/> sennosides-docusate sodium (SENOKOT-S) 8.6-50 mg per tablet	1 tablet, oral, 2 times daily, Post-op Hold for diarrhea.
<input type="checkbox"/> bisacodyl (DULCOLAX) suppository	10 mg, rectal, daily PRN, constipation, (if with persistent constipation), Post-op
<input type="checkbox"/> polyethylene glycol (MIRALAX) packet	17 g, oral, daily PRN, constipation, Post-op
<input type="checkbox"/> docusate sodium (COLACE) capsule	100 mg, oral, daily PRN, constipation, Post-op
<input type="checkbox"/> Other	

Antiemetics - HHM, HMSJ, HMW, HMSTC Only

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

<input type="checkbox"/>	promethazine (PHENERGAN) tablet	12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is able to tolerate oral medication.
<input type="checkbox"/>	promethazine (PHENERGAN) suppository	12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication.
<input type="checkbox"/>	Other	

Antiemetics - HMSTJ Only

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

Antiemetics - HMSL, HMWB Only

<input type="checkbox"/>	ondansetron (ZOFTRAN) IV or Oral	"Or" Linked Panel
<input type="checkbox"/>	ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet	4 mg, oral, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is able to tolerate oral medication.
<input type="checkbox"/>	ondansetron (ZOFTRAN) 4 mg/2 mL injection	4 mg, intravenous, every 8 hours PRN, nausea, vomiting, Post-op Give if patient is UNable to tolerate oral medication OR if a faster onset of action is required.
<input type="checkbox"/>	promethazine (PHENERGAN) IV or Oral or Rectal	"Or" Linked Panel
<input type="checkbox"/>	promethazine (PHENERGAN) 12.5 mg in sodium chloride 0.9 % 0.9 % 20 mL for Alaris pump syringe option	12.5 mg, intravenous, at 60 mL/hr, for 20 Minutes, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required.
<input type="checkbox"/>	promethazine (PHENERGAN) tablet	12.5 mg, oral, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is able to tolerate oral medication.
<input type="checkbox"/>	promethazine (PHENERGAN) suppository	12.5 mg, rectal, every 6 hours PRN, nausea, vomiting, Post-op Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication.

Other

Respiratory Medications

<input type="checkbox"/> albuterol (PROVENTIL) nebulizer solution	2.5 mg, nebulization, Respiratory Therapy - every 4 hours, For 7 Days, Post-op Aerosol Delivery Device: if (answer = Intrapulmonary Percussive Ventilation (Meta-Neb Device)) Meta-Neb Indications:
<input type="checkbox"/> ipratropium (ATROVENT) 0.02 % nebulizer solution	0.5 mg, nebulization, Respiratory Therapy - every 4 hours, For 7 Days, Post-op Aerosol Delivery Device: if (answer = Intrapulmonary Percussive Ventilation (Meta-Neb Device)) Meta-Neb Indications:
<input type="checkbox"/> acetylcysteine 200 mg/mL (20 %) inhalation dose	400 mg, nebulization, Respiratory Therapy - every 12 hours, For 7 Days, Post-op Aerosol Delivery Device: Intrapulmonary Percussive Ventilation (Meta-Neb Device) if (answer = Intrapulmonary Percussive Ventilation (Meta-Neb Device)) Meta-Neb Indications: Meta-Neb Indications: Inadequate Secretion Clearance

Other

Other Medications

<input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet - POD #5	81 mg, oral, daily, Starting S+5, Post-op
<input type="checkbox"/> magnesium oxide (MAG-OX) tablet - POD #5	400 mg, oral, 2 times daily, Starting S+5, Post-op
<input type="checkbox"/> multivitamin with minerals tablet - POD #5	1 tablet, oral, daily, Starting S+5, Post-op
<input type="checkbox"/> calcium carbonate-vitamin D3 500 mg-200 unit per tablet - POD #5	1 tablet, oral, 2 times daily, Starting S+5, Post-op
<input type="checkbox"/> ergocalciferol (VITAMIN D2) capsule - POD #5	50,000 Units, oral, weekly, Starting S+5, Post-op
<input type="checkbox"/> Other	

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

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

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

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

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

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

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

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

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

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

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

<input type="radio"/> Low Risk of DVT	
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<input type="checkbox"/> Low Risk (Single Response)	
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<input type="radio"/> Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation PACU & Post-op
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<input type="radio"/> Moderate Risk of DVT - Surgical	
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Address pharmacologic prophylaxis by selecting one of the following. Mechanical prophylaxis is optional unless pharmacologic prophylaxis is contraindicated.

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

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

<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 (TIME CRITICAL), PACU & Post-op Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous, PACU & Post-op
<input type="checkbox"/> High Risk of DVT - Surgical	
Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.	
<input type="checkbox"/> High Risk	
<input type="checkbox"/> High risk of VTE	Routine, Once, PACU & Post-op
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response)	
<input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: if (answer = Other) Other anticoagulant therapy: PACU & Post-op
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1 For Patients with CrCL LESS than 30 mL/min
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min

() 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: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
[] 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 ordering 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	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: if (answer = Other) Other anticoagulant therapy: 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)	

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

() 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: if (answer = Other) Other anticoagulant therapy: 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: if (answer = Other (Please specify)) Specify Other Indication:
() 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)	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1
() enoxaparin (LOVENOX) syringe	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 > 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: if (answer = Other (Please specify)) Specify Other Indication:

warfarin (COUMADIN) tablet oral, daily at 1700 (TIME CRITICAL), Starting S+1, PACU & Post-op
Indication:
if (answer = Other (Specify indication & Target INR))
Specify indication & Target INR (free text):
if (answer = LVAD (Specify Target INR))
Target INR:

Pharmacy consult to manage warfarin (COUMADIN) STAT, Until discontinued, Starting S
Indication:
if (answer = Other (Specify indication & Target INR))
Specify indication & Target INR (free text):
if (answer = LVAD (Specify Target INR))
Target INR:

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

Other

DVT Risk and Prophylaxis Tool (Single Response)

Low Risk Definition Moderate Risk Definition

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

Both pharmacologic AND mechanical prophylaxis must be addressed.

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

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

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

Central line Acute spinal cord injury with paresis

History of DVT or family history of VTE Multiple major traumas

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

Less than fully and independently ambulatory Acute ischemic stroke

Estrogen therapy History of PE

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Low Risk of DVT

Low Risk (Single Response)

Low risk of VTE Routine, Once
Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation
PACU & Post-op

Moderate Risk of DVT - Surgical

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

<input type="checkbox"/>	Moderate Risk	
<input type="checkbox"/>	Moderate risk of VTE	Routine, Once, PACU & Post-op
<input type="checkbox"/>	Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response)	
<input type="checkbox"/>	Patient is currently receiving therapeutic anticoagulation	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: if (answer = Other) Other anticoagulant therapy: PACU & Post-op
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/>	enoxaparin (LOVENOX) injection (Single Response)	
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1 For Patients with CrCL LESS than 30 mL/min
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe - For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min
<input type="checkbox"/>	fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1, PACU & Post-op If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT):
<input type="checkbox"/>	heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op
<input type="checkbox"/>	heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs)	5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/>	warfarin (COUMADIN) tablet	oral, daily at 1700 (TIME CRITICAL), Starting S+1, PACU & Post-op Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:

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

<input type="checkbox"/> heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs)	5,000 Units, subcutaneous, every 12 hours, PACU & Post-op Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 (TIME CRITICAL), PACU & Post-op Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous, PACU & Post-op
<input type="checkbox"/> High Risk of DVT - Surgical	
Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.	
<input type="checkbox"/> High Risk	
<input type="checkbox"/> High risk of VTE	Routine, Once, PACU & Post-op
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response)	
<input type="checkbox"/> Patient is currently receiving therapeutic anticoagulation	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: if (answer = Other) Other anticoagulant therapy: PACU & Post-op
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1 For Patients with CrCL LESS than 30 mL/min
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min

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

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

[] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty)
Surgical Patient (Single Response)

<p>() Patient is currently receiving therapeutic anticoagulation</p>	<p>Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following: if (answer = Other) Other anticoagulant therapy: PACU & Post-op</p>
<p>() Contraindications exist for pharmacologic prophylaxis</p>	<p>Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): PACU & Post-op</p>
<p>() apixaban (ELIQUIS) tablet</p>	<p>2.5 mg, oral, every 12 hours, Starting S+1, PACU & Post-op Indications: if (answer = Other (Please specify)) Specify Other Indication:</p>
<p>() aspirin chewable tablet</p>	<p>162 mg, oral, daily, Starting S+1, PACU & Post-op</p>
<p>() aspirin (ECOTRIN) enteric coated tablet</p>	<p>162 mg, oral, daily, Starting S+1, PACU & Post-op</p>
<p>() enoxaparin (LOVENOX) injection (Single Response)</p>	
<p>() enoxaparin (LOVENOX) syringe</p>	<p>40 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1</p>
<p>() enoxaparin (LOVENOX) syringe</p>	<p>30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1</p>
<p>() enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min</p>	<p>30 mg, subcutaneous, daily at 0600 (TIME CRITICAL), Starting S+1 For Patients with CrCL LESS than 30 mL/min.</p>
<p>() enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min</p>	<p>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.</p>
<p>() enoxaparin (LOVENOX) syringe - For Patients weight between 140 kg or GREATER and CrCl GREATER than 30 mL/min</p>	<p>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</p>
<p>() fondaparinux (ARIXTRA) injection</p>	<p>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):</p>
<p>() heparin (porcine) injection</p>	<p>5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM, PACU & Post-op</p>
<p>() heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs)</p>	<p>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.</p>

<input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission	10 mg, oral, daily at 0600 (TIME CRITICAL), Starting S+1, PACU & Post-op To be Given on Post Op Day 1. Indications: if (answer = Other (Please specify)) Specify Other Indication:
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 (TIME CRITICAL), Starting S+1, PACU & Post-op Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication: if (answer = Other (Specify indication & Target INR)) Specify indication & Target INR (free text): if (answer = LVAD (Specify Target INR)) Target INR:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s): PACU & Post-op
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous, PACU & Post-op

Other

Labs

Laboratory STAT Upon Arrival

<input type="checkbox"/> Lactic acid level	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> CBC with platelet and differential	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> Comprehensive metabolic panel	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> Ionized calcium	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> Magnesium level	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> Phosphorus level	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> LDH	STAT For 1 Occurrences, Post-op
<input type="checkbox"/> Vitamin D 25 hydroxy level	Once For 1 Occurrences, Post-op
<input type="checkbox"/> Other	

Labs every 6 hours x 3

<input type="checkbox"/> Hemoglobin and hematocrit	Every 6 hours For 3 Occurrences 6 hours after Arrival, Post-op
<input type="checkbox"/> Basic metabolic panel	Every 6 hours For 3 Occurrences 6 hours after Arrival, Post-op
<input type="checkbox"/> Other	

Troponin x 3

<input type="checkbox"/>	Troponin	Every 8 hours For 3 Occurrences
<input type="checkbox"/>	Other	

Laboratory Every Morning x 3 days

<input type="checkbox"/>	CBC with platelet and differential	AM draw repeats For 7 Days, Post-op
<input type="checkbox"/>	Comprehensive metabolic panel	AM draw repeats For 3 Days, Post-op
<input type="checkbox"/>	Magnesium level	AM draw repeats For 7 Days, Post-op
<input type="checkbox"/>	Phosphorus level	AM draw repeats For 7 Days, Post-op
<input type="checkbox"/>	Ionized calcium	AM draw repeats For 3 Days, Post-op
<input type="checkbox"/>	LDH	AM draw repeats For 3 Days, Post-op
<input type="checkbox"/>	Other	

Immunosupression Levels

<input type="checkbox"/>	FK506 Tacrolimus level, trough	AM draw repeats, Starting S+1 For 7 Days, Post-op
<input type="checkbox"/>	Other	

Post Transplant Labs Mondays x 3

<input type="checkbox"/>	Cytomegalovirus by PCR	Every Monday For 3 Occurrences Specimen Source: Post-op
<input type="checkbox"/>	Other	

Arterial Blood Gas

<input type="checkbox"/>	Arterial blood gas	STAT For 1 Occurrences, Post-op
<input type="checkbox"/>	Arterial blood gas	Every 6 hours For 3 Occurrences Every 6 hours x 3, Post-op
<input type="checkbox"/>	Arterial blood gas	AM draw repeats For 3 Days, Post-op
<input type="checkbox"/>	Other	

Microbiology

<input type="checkbox"/>	Blood culture x 2	"And" Linked Panel
<input type="checkbox"/>	Blood Culture (Aerobic & Anaerobic)	Once, Starting S For 1 Occurrences, Blood Upon arrival to the unit
		Collect before antibiotics given. Blood cultures should be ordered x2, with each set drawn from a different peripheral site. If unable to draw both sets from a peripheral site, one set may be drawn from a central line; an IV line should NEVER be used., Post-op

<input type="checkbox"/> Blood Culture (Aerobic & Anaerobic)	<p>Once For 1 Occurrences, Blood Upon arrival to the unit.</p> <p>Collect before antibiotics given. Blood cultures should be ordered x2, with each set drawn from a different peripheral site. If unable to draw both sets from a peripheral site, one set may be drawn from a central line; an IV line should NEVER be used., Post-op</p>
<input type="checkbox"/> Blood culture x 2	<p>"And" Linked Panel</p>
<input type="checkbox"/> Blood Culture (Aerobic & Anaerobic)	<p>Conditional Frequency, Blood One activation if temperature greater than 99.9 Fahrenheit.</p> <p>Collect before antibiotics given. Blood cultures should be ordered x2, with each set drawn from a different peripheral site. If unable to draw both sets from a peripheral site, one set may be drawn from a central line; an IV line should NEVER be used., Post-op</p>
<input type="checkbox"/> Blood Culture (Aerobic & Anaerobic)	<p>Conditional Frequency, Blood One activation if temperature greater than 99.9 Fahrenheit.</p> <p>Collect before antibiotics given. Blood cultures should be ordered x2, with each set drawn from a different peripheral site. If unable to draw both sets from a peripheral site, one set may be drawn from a central line; an IV line should NEVER be used., Post-op</p>
<input type="checkbox"/> Urinalysis screen and microscopy, with reflex to culture	<p>Conditional Frequency Specimen Source: Urine Specimen Site: One activation if temperature greater than 99.9 Fahrenheit., Post-op</p>
<input type="checkbox"/> Other	

Cardiology

Cardiology

<input type="checkbox"/> ECG 12 lead	<p>STAT, Once For 1 Occurrences Clinical Indications: Post-Op Surgery if (answer = Other:) Other: Interpreting Physician: STAT upon arrival to unit, Post-op</p>
<input type="checkbox"/> ECG 12 lead	<p>Routine, Daily, Starting S+1 For 3 Days Clinical Indications: Post-Op Surgery if (answer = Other:) Other: Interpreting Physician: Every morning times 3 days, Post-op</p>
<input type="checkbox"/> Other	

Imaging

X-Ray

<input type="checkbox"/> XR Chest 1 Vw Portable	STAT, 1 time imaging For 1 Occurrences STAT upon arrival to unit., Post-op
<input type="checkbox"/> XR Chest 1 Vw Portable	Routine, Daily imaging, Starting S+1 For 7 Days AM Every morning x 7 days, Post-op
<input type="checkbox"/> XR Chest 1 Vw Portable	STAT, Conditional Frequency For 1 Unlimited activations if temperature greater than 99.9 degrees Fahrenheit., Post-op
<input type="checkbox"/> Other	

Respiratory

Respiratory Therapy

<input type="checkbox"/> Suctioning	Routine, Every 4 hours Route: Endotracheal if (answer = Other (Specify)) Specify: Bag and suction with coude catheter only. Do not suction if PEEP is more than 10, unless absolutely necessary, Post-op
<input type="checkbox"/> Incentive spirometry	Routine, Every hour Start when extubated., Post-op
<input type="checkbox"/> Encourage deep breathing and coughing	Routine, Every 2 hours Start when extubated., Post-op
<input type="checkbox"/> Other	

Consults

For Physician Consult orders use sidebar

Consults

<input type="checkbox"/> Consult Diabetes/Endocrinology	Reason for Consult? Patient/Clinical information communicated? if (answer = Answering service) Additional information: Patient/clinical information communicated? if (answer = Consultant not contacted) Will you contact the consultant? if (answer = Answering service notified) Additional information:
<input type="checkbox"/> Consult to PT eval and treat	Special Instructions: Weight Bearing Status:
<input type="checkbox"/> Consult to OT eval and treat	Special Instructions: Weight Bearing Status:

[] Consult to Nutrition Services	Reason For Consult? Other (Specify) if (answer = Other (Specify)) Specify: Specify: Post Transplant Diet Education Post-op, Registered Dietitian
[] Consult Methodist Rehab Associates	Reason for Consult: PM&R Evaluation if (answer = Other) Specify: Post-op
[] Consult to Case Management	Consult Reason: Other specify if (answer = Other specify) Specify: if (answer = Home Health) Face-to-Face Date: Reasons for Home Health Care: Home Health Services: if (answer = Skilled Nursing Evaluation & Treatment) Times per week: For: Days/Week/Weeks: if (answer = Physical Therapy Evaluation & Treatment) (PT) Times per week: For: Days/Week/Weeks: if (answer = Occupational Therapy Evaluation & Treatment) Times per week: For: Days/Week/Weeks: if (answer = Speech Language Pathology Evaluation & Treatment) Times per week: For: Days/Week/Weeks: if (answer = Social Worker) Times per week: For: Days/Week/Weeks: if (answer = Home Health Aide) Times per week: For: Days/Week/Weeks: if (answer = Home Infusion) IV infusion needs: if (answer = Labs) IV Infusion Labs: Every: Lab results called to: if (answer = IV Fluids) Solution:

How often:
Start date:
Stop date:
if (answer = Antibiotics)
Antibiotic(s), please list:
Start date:
Stop date:
if (answer = Nutritional Supplies)
Nutritional DME:
if (answer = Bolus feeding)
Rate:
Formula:
if (answer = Continuous feeding)
Rate:
Formula:
if (answer = Home Wound Care)
Wound care questions:
if (answer = Dressing Instructions)
How often:
Clean with:
Cover with:
Duration:
if (answer = Pleurx)
PleurX choices:
Change every:
PleurX Duration:
if (answer = Wound vac)
Change how often:
Pressure (mmHg):
Therapy Settings:
if (answer = Other)
Specify:
if (answer = Dynamic Pressure Control)
DCP Ratio:
Intensity:
Foam Type:
Type of Wound:
if (answer = Other)
Specify:
if (answer = Ostomy supplies)
Special ostomy supplies:
Clinical Findings:
if (answer = Other:)
Other Clinical Findings:
Homebound Status:
if (answer = Other:)
Other Homebound Status:
if (answer = Leaving home is medically contraindicated due to)
Contraindication:

Special Instructions:
Resume home health services with previous home health agency prior to the hospital admission:
Face to Face Cert Statement:
if (answer = DME)
DME Diagnosis:
Type of DME:
if (answer = Mobility Aids)
MOBILITY AIDS: Per Payer requirements; only ONE Mobility Aid may be chosen from this list:
if (answer = Walkers (With 5 inch Wheels))
Walkers (With 5 inch wheels):
if (answer = Walkers (Without Wheels))
Walkers (Without Wheels):
if (answer = Wheelchair)
Wheelchair:
if (answer = Canes)
Canes:
if (answer = Crutches)
Crutches:
if (answer = 3 in 1 Bedside Commode)
3-in-1 Bedside Commode:
if (answer = Respiratory Equipment)
Oxygen:
if (answer = O2 Portable Gas)
Continuous or PRN Oxygen:
O2 Duration:
O2 Sat on Room Air, at Rest %:
O2 Sat on Room Air, During Exertion %:
O2 Sat on Oxygen with Exertion % demonstrates improvement (above 88%):
O2 Device:
O2 Flowrate (L/Min) Setting:
INDICATIONS for Ordering Oxygen: Must enter Lung Disease or Hypoxia Related Symptoms:
if (answer = Lung Disease Diagnosis)
INDICATIONS for Ordering Oxygen: Must enter Lung Disease Diagnosis or Hypoxia Related Symptoms - Lung Disease Diagnosis:
if (answer = Hypoxia Related Symptoms)
Hypoxia Related Symptoms:
if (answer = Nebulizer)
Nebulizer Med:
if (answer = Albuterol)
Albuterol dose:
if (answer = Xopenex)
Xopenex dose:
if (answer = Mucomyst)
Mucomyst dose:
if (answer = Atrovent)

Atrovent dose:
INDICATIONS for Ordering Nebulizer: Must enter Lung Disease or Hypoxia Related Symptoms:
if (answer = Lung Disease Diagnosis)
INDICATIONS for Ordering Nebulizer: Must enter Lung Disease Diagnosis or Hypoxia Related Symptoms - Lung Disease Diagnosis:
if (answer = Hypoxia Related Symptoms)
Hypoxia Related Symptoms:
if (answer = Trach supplies)
Type:
Size of tube:
if (answer = Home ventilator)
Home ventilator settings:
if (answer = CPAP)
Pressure:
if (answer = BIPAP)
IPAP:
EPAP:
if (answer = O2 Bleed in Rate)
Liter flow:
if (answer = Portable O2 Generator)
Continuous or PRN Oxygen:
O2 Duration:
O2 Sat on Room Air, at Rest %:
O2 Sat on Room Air, During Exertion %:
O2 Sat on Oxygen with Exertion % demonstrates improvement (above 88%):
O2 Device:
O2 Flowrate (L/Min) Setting:
INDICATIONS for Ordering Oxygen: Must enter Lung Disease or Hypoxia Related Symptoms:
if (answer = Lung Disease Diagnosis)
INDICATIONS for Ordering Oxygen: Must enter Lung Disease Diagnosis or Hypoxia Related Symptoms - Lung Disease Diagnosis:
if (answer = Hypoxia Related Symptoms)
Hypoxia Related Symptoms:
if (answer = Hospital Bed)
Hospital Bed:
if (answer = Gel Overlay)
Indicate which of the following conditions describe the patient. Answer all that apply:
if (answer = Alternating Pressure Mattress)
Indicate which of the following conditions describe the patient. Answer all that apply:
if (answer = Low Air Loss Mattress)
Additional Medical Information - select all that apply:
if (answer = Semi-Electric Hospital Bed with Split Siderails)
Pressure ulcer - check all that apply:
if (answer = Semi-Electric Hospital Bed with Full Rails)

Pressure ulcer - check all that apply:
if (answer = Other Equipment (specify))
Other Equipment:
if (answer = Other (specify))
Other:
if (answer = Diabetic supplies)
Diabetic supplies:
Face-to-Face Date:
Clinical Findings:
if (answer = Other:)
Other Clinical Findings:
Special Instructions:
Specify: Lung Transplant; arrange home nebulizer machine
Post-op

Consult to Transplant Social Work

Reason for Consult?
Organ Transplant: Lung
Post-op

Consult to Speech Language Pathology

Routine, Once, Starting S+1
Reason for consult:
if (answer = Other)
Reason for SLP?
Post-op

Other