

General

Common Present on Admission Diagnosis

<input type="checkbox"/> Present on Admission	Details
<input type="checkbox"/> Abdominal pain	Details
<input type="checkbox"/> Back pain	Details
<input type="checkbox"/> Chest pain	Details
<input type="checkbox"/> Cough	Details
<input type="checkbox"/> COVID - 19	Details
<input type="checkbox"/> Dizziness	Details
<input type="checkbox"/> Fall	Details
<input type="checkbox"/> Fever	Details
<input type="checkbox"/> Headache	Details
<input type="checkbox"/> Hypertension	Details
<input type="checkbox"/> Nausea	Details
<input type="checkbox"/> Shortness of breath	Details
<input type="checkbox"/> Vomiting	Details
<input type="checkbox"/> Weakness-generalized	Details

Admission or Observation (Single Response) (Selection Required)

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

Admission or Observation (Single Response)

Patient has active status order on file

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

Admission (Single Response)

Patient has active status order on file.

Admit to inpatient

Admitting Physician:
 Level of Care:
 Patient Condition:
 Bed request comments:
 Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and progress notes, I expect that the patient will need hospital services for two or more midnights.

Code Status (Single Response)

Full code Code Status decision reached by:

DNR (Selection Required)

DNR (Do Not Resuscitate) Did the patient/surrogate require the use of an interpreter?
 Did the patient/surrogate require the use of an interpreter?
 Does patient have decision-making capacity?

Consult to Palliative Care Service Priority:
 Reason for Consult?
 Order?
 Name of referring provider:
 Enter call back number:

Consult to Social Work Reason for Consult:

Modified Code Did the patient/surrogate require the use of an interpreter?
 Did the patient/surrogate require the use of an interpreter?
 Does patient have decision-making capacity?
 Modified Code restrictions:

Treatment Restrictions I understand that if the patient is NOT in a cardiopulmonary arrest, the selected treatments will NOT be provided. I understand that all other unselected medically indicated treatments will be provided.
 Treatment Restriction decision reached by:
 Specify Treatment Restrictions:

Isolation

Airborne isolation status

Airborne isolation status Details

Mycobacterium tuberculosis by PCR - If you suspect Tuberculosis, please order this test for rapid diagnostics. Once, Sputum

Contact isolation status Details

Droplet isolation status Details

Enteric isolation status Details

Precautions

Aspiration precautions Details

Fall precautions Increased observation level needed:

Latex precautions Details

Seizure precautions Increased observation level needed:

Nursing

Vital Signs

Vital signs - Per Unit Protocol Routine, Per unit protocol

Vital signs-Q4H Routine, Every 4 hours For Until specified

Telemetry Order

Telemetry **"And" Linked Panel**

<input type="checkbox"/> Telemetry monitoring	Routine, Continuous Order: Place in Centralized Telemetry Monitor: EKG Monitoring Only (Telemetry Box) Reason for telemetry: Chest pain syndrome Can be off of Telemetry for tests and baths? Yes
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<input type="checkbox"/> Telemetry Additional Setup Information	Routine, Continuous High Heart Rate (BPM): 120 Low Heart Rate(BPM): 50 High PVC's (per minute): 10 High SBP(mmHg): 175 Low SBP(mmHg): 100 High DBP(mmHg): 95 Low DBP(mmHg): 40 Low Mean BP: 60 High Mean BP: 120 Low SPO2(%): 94
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Activity

<input type="checkbox"/> Strict bed rest	Routine, Until discontinued, Starting S
<input type="checkbox"/> Bed rest with bathroom privileges	Routine, Until discontinued, Starting S Bathroom Privileges: with bathroom privileges
<input type="checkbox"/> Ambulate with assistance	Routine, 3 times daily Specify: with assistance
<input type="checkbox"/> Activity as tolerated	Routine, Until discontinued, Starting S Specify: Activity as tolerated

Nursing

<input checked="" type="checkbox"/> Head of bed 30 degrees	Routine, Until discontinued, Starting S Head of bed: 30 degrees
<input checked="" type="checkbox"/> Daily weights	Routine, Daily
<input checked="" type="checkbox"/> Intake and Output	Routine, Every 8 hours Including bowel movements
<input type="checkbox"/> Bedside glucose - every 4 hours	Routine, Every 4 hours
<input type="checkbox"/> Bedside glucose - AC & HS	Routine, 4 times daily 0-30 minutes before meals and at bedtime
<input type="checkbox"/> Nasogastric Tube Orders	
<input type="checkbox"/> Nasogastric tube insertion	Routine, Once Type:
<input type="checkbox"/> Nasogastric tube maintenance	Routine, Until discontinued, Starting S Tube Care Orders:
<input type="checkbox"/> Insert and Maintain Foley	
<input type="checkbox"/> Insert Foley catheter	Routine, Once Type: Size: Urinometer needed:
<input type="checkbox"/> Foley Catheter Care	Routine, Until discontinued, Starting S Orders: Maintain
<input type="checkbox"/> Change foley catheter	Routine, Once
<input type="checkbox"/> Oral care	Routine, Every 8 hours

Notify

<input checked="" type="checkbox"/> Notify Physician (Specify)	Routine, Until discontinued, Starting S, Active bleeding
<input checked="" type="checkbox"/> Notify Physician (Specify)	Routine, Until discontinued, Starting S, Change in condition or Glasgow Coma Score less than 13

<input checked="" type="checkbox"/> Notify Physician(vitals,output,pulse ox)	Routine, Until discontinued, Starting S Temperature greater than: 101.5 Temperature less than: Systolic BP greater than: 160 Systolic BP less than: 80 Diastolic BP greater than: Diastolic BP less than: Heart rate greater than (BPM): 110 Heart rate less than (BPM): 50 Respiratory rate greater than: 30 Respiratory rate less than: 10 SpO2 less than: Urine Output less than: 30ml/hr or less than 250ml/8 hours Output (Specify) greater than: Other:
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Diet

<input type="checkbox"/> NPO	Diet effective now, Starting S NPO: Pre-Operative fasting options:
<input type="checkbox"/> NPO-Except ice chips	Diet effective now, Starting S NPO: Except Ice chips Pre-Operative fasting options:
<input type="checkbox"/> Diet-500ml fluid restriction	Diet effective now, Starting S Diet(s): Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Fluid Restriction 500 ml Foods to Avoid:
<input type="checkbox"/> Diet-1000ml fluid restriction	Diet effective now, Starting S Diet(s): Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Fluid Restriction 1000 ml Foods to Avoid:
<input type="checkbox"/> Diet-2gm Sodium	Diet effective now, Starting S Diet(s): 2 GM Potassium Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid:
<input type="checkbox"/> Diet-1800 Carb Control Diabetic	Diet effective now, Starting S Diet(s): Other Diabetic/Cal Diabetic/Calorie: 1800 Kcal/202 gm Carbohydrate Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid:
<input type="checkbox"/> Diet-Renal(80GM, 2-3GM Na, 2-3GM K)	Diet effective now, Starting S Diet(s): Renal (80GM Pro, 2-3GM Na, 2-3GM K) Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid:
<input type="checkbox"/> Diet-Clear liquids	Diet effective now, Starting S Diet(s): Clear Liquids Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid:

<input type="checkbox"/> Diet-Full liquids	Diet effective now, Starting S Diet(s): Full Liquids Advance Diet as Tolerated? IDDSI Liquid Consistency: Fluid Restriction: Foods to Avoid:
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IV Fluids

Peripheral IV Access

<input checked="" type="checkbox"/> Initiate and maintain IV	
<input checked="" type="checkbox"/> Insert peripheral IV	Routine, Once
<input checked="" type="checkbox"/> sodium chloride 0.9 % flush	10 mL, intravenous, every 12 hours scheduled
<input checked="" type="checkbox"/> sodium chloride 0.9 % flush	10 mL, intravenous, PRN, line care

IV Fluid

<input type="checkbox"/> sodium chloride 0.45 % infusion 1000 mL	intravenous, at 75 mL/hr, continuous
<input type="checkbox"/> sodium chloride 0.45 % with potassium chloride 20 mEq/L infusion	75 mL/hr, intravenous, continuous
<input type="checkbox"/> dextrose 5%-0.45% sodium chloride 1,000 mL infusion	75 mL/hr, intravenous, continuous

Medications

Pharmacy Consults

<input checked="" type="checkbox"/> Pharmacy consult to manage dosing of medication	Routine, Until discontinued, Starting S Adjust dose for: renal function
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Medications

<input type="checkbox"/> zinc sulfate (ZINCATE) capsule	1 capsule, oral, daily
<input type="checkbox"/> magnesium oxide tablet	400 mg, oral, 3 times daily
<input type="checkbox"/> magnesium sulfate 2 g in sodium chloride 0.45 % 0.45 % 100 mL IVPB	2 g, intravenous, for 1 Hours, once, For 1 Doses
<input type="checkbox"/> octreotide (SANDOSTATIN) IV Bolus Once AND Maintenance 25 mcg/hr	"And" Linked Panel
<input type="checkbox"/> octreotide (SANDOSTATIN) bolus injection	100 mcg, intravenous, once, For 1 Doses Bolus once initial dose. Infusion to start immediately after bolus. May cause Q-T interval prolongation.
<input type="checkbox"/> octreotide (SandoSTATIN) maintenance infusion	25 mcg/hr, intravenous, for 40 Hours, continuous May cause Q-T interval prolongation
<input type="checkbox"/> octreotide (SANDOSTATIN) IV Bolus Once AND Maintenance 50 mcg/hr	"And" Linked Panel
<input type="checkbox"/> octreotide (SANDOSTATIN) bolus injection	100 mcg, intravenous, once, For 1 Doses Bolus once initial dose. Infusion to start immediately after bolus. May cause Q-T interval prolongation.
<input type="checkbox"/> octreotide (SandoSTATIN) maintenance infusion	50 mcg/hr, intravenous, continuous May cause Q-T interval prolongation
<input type="checkbox"/> pantoprazole (PROTONIX) IV or ORAL	"Or" Linked Panel
<input type="checkbox"/> pantoprazole (PROTONIX) EC tablet	40 mg, oral, daily at 0600 Indication(s) for Proton Pump Inhibitor (PPI) Therapy:
<input type="checkbox"/> pantoprazole (PROTONIX) 40 mg in sodium chloride 0.9 % 10 mL injection	40 mg, intravenous, daily at 0600 Indication(s) for Proton Pump Inhibitor (PPI) Therapy:

Ascites

<input type="checkbox"/> furosemide (LASIX) injection	40 mg, intravenous, 2 times daily at 0900, 1700
<input type="checkbox"/> furosemide (LASIX) tablet	40 mg, oral, 2 times daily at 0900, 1700
<input type="checkbox"/> spironolactone (ALDACTONE) tablet	100 mg, oral, daily MONITOR POTASSIUM LEVELS. AVOID SALT SUBSTITUTES UNLESS APPROVED BY MD.
<input type="checkbox"/> bumetanide (BUMEX) injection	1 mg, intravenous, daily
<input type="checkbox"/> bumetanide (BUMEX) tablet	1 mg, oral, daily

<input type="checkbox"/>	hydrochlorothiazide (HYDRODIURIL)	25 mg, oral, daily
<input type="checkbox"/>	albumin human 25 % bottle	50 mL, intravenous, at 50 mL/hr, for 60 Minutes, once, For 1 Doses Indication:

Hepatorenal Syndrome

<input type="checkbox"/>	midodrine (PROAMATINE) tablet	10 mg, oral, 3 times daily at 0900, 1300, 1700 BP HOLD parameters for this order:
<input type="checkbox"/>	octreotide (SANDOSTATIN) injection	100 mcg, subcutaneous, every 8 hours
<input type="checkbox"/>	albumin human 25 % bottle	50 mL, intravenous, at 50 mL/hr, for 60 Minutes, every 8 hours Indication:
<input type="checkbox"/>	albumin human 5 % bottle	12.5 g, intravenous, at 125 mL/hr, for 120 Minutes, every 4 hours, For 2 Doses Indication:
<input type="checkbox"/>	albumin human 5 % bottle	25 g, intravenous, at 250 mL/hr, for 120 Minutes, every 4 hours, For 2 Doses Indication:

Spont. Bacterial Peritonitis

<input type="checkbox"/>	Community Acquired SBP - Ceftriaxone	"And" Linked Panel
<input type="checkbox"/>	cefTRIAxone (ROCEPHIN) IV	2 g, intravenous, for 30 Minutes, every 24 hours Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	Hospital Acquired SBP (Hospitalization within the past 90 days OR outpatient intravenous therapy within the past 30 days) - Cefepime PLUS Metronidazole PLUS Vancomycin	
<input type="checkbox"/>	ceFEPime (MAXIPIME) IV	2 g, intravenous, every 8 hours Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	metronidazole (FLAGYL)	500 mg, intravenous, for 30 Minutes, every 8 hours Per Med Staff Policy, R.Ph. will automatically switch IV to equivalent PO dose when above approved criteria are satisfied: Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required)	
<input type="checkbox"/>	vancomycin (VANCOGIN)	15 mg/kg, intravenous, every 12 hours On call to cath lab. Transport antibiotics with patient to cath lab and administer prior to start of procedure Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	Pharmacy consult to manage vancomycin	STAT, Until discontinued, Starting S Indication: Reason for Therapy: Bacterial Infection Suspected Indication: Duration of Therapy (Days):
<input type="checkbox"/>	Severe Penicillin Allergy - Aztreonam PLUS Vancomycin +/- Metronidazole	
<input type="checkbox"/>	aztreonam (AZACTAM) IV	2 g, intravenous, every 8 hours Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	metronidazole (FLAGYL)	500 mg, intravenous, every 8 hours Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	vancomycin 15 mg/kg IV + Pharmacy Consult (Selection Required)	
<input type="checkbox"/>	vancomycin (VANCOGIN)	15 mg/kg, intravenous, every 12 hours Reason for Therapy: Bacterial Infection Suspected Indication:
<input type="checkbox"/>	Pharmacy consult to manage vancomycin	STAT, Until discontinued, Starting S Indication:

<input type="checkbox"/> albumin human 25 % bottle	50 mL, intravenous, at 50 mL/hr, once, For 1 Doses Indication:
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Encephalopathy

<input type="checkbox"/> lactulose (CHRONULAC) 10 gram/15 mL solution	10 g, oral, every 1 hour STOP AFTER FIRST BOWEL MOVEMENT AND START 3 TIMES DAILY DOSING.
<input type="checkbox"/> lactulose (CHRONULAC) 10 gram/15 mL solution	10 g, oral, 3 times daily START AFTER FIRST BOWEL MOVEMENT. HOLD FOR MORE THAN 5 BOWEL MOVEMENTS IN 24 HOURS.
<input type="checkbox"/> lactulose solution (Enema)	200 g, rectal, once, For 1 Doses
<input type="checkbox"/> neomycin (MYCIFRADIN) tablet	1,000 mg, oral, 3 times daily Reason for Therapy:
<input type="checkbox"/> rifaximin (XIFAXAN) tablet	550 mg, oral, 2 times daily Indication:

Variceal Bleeding / Portal HTN

<input type="checkbox"/> propranolol (INDERAL) tablet	20 mg, oral, 2 times daily at 0600, 1800 BP & HR HOLD parameters for this order: Contact Physician if:
<input type="checkbox"/> carvedilol (COREG) tablet	6.25 mg, oral, 2 times daily at 0600, 1800 BP & HR HOLD parameters for this order: Contact Physician if:
<input type="checkbox"/> phytonadione (AQUA-MEPHYTON) injection	10 mg, subcutaneous, daily Indication:

PRN Mild Pain (Pain score 1-3)

(adjust dose for renal/liver function and age)

<input checked="" type="checkbox"/> acetaminophen (TYLENOL) tablet OR oral solution	"Or" Linked Panel Maximum of 4 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources)
<input checked="" type="checkbox"/> acetaminophen (TYLENOL) tablet	650 mg, oral, every 6 hours PRN, mild pain (score 1-3) Give if patient able to take oral tablet medication.
<input checked="" type="checkbox"/> acetaminophen (TYLENOL)suspension	650 mg, oral, every 6 hours PRN, mild pain (score 1-3) Give if patient cannot receive oral tablet but can receive oral solution.

Antiemetics

<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral (Selection Required)	"Or" Linked Panel
<input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet	4 mg, oral, every 8 hours PRN, nausea, vomiting Give if patient is able to tolerate oral medication.
<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection	4 mg, intravenous, every 8 hours PRN, nausea, vomiting 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 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 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 Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication.

Antiemetics

<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral (Selection Required)	"Or" Linked Panel
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<input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet	4 mg, oral, every 8 hours PRN, nausea, vomiting Give if patient is able to tolerate oral medication.
<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection	4 mg, intravenous, every 8 hours PRN, nausea, vomiting 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 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 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 Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication.

Antiemetics

<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) IV or Oral (Selection Required)	"Or" Linked Panel
<input checked="" type="checkbox"/> ondansetron ODT (ZOFTRAN-ODT) disintegrating tablet	4 mg, oral, every 8 hours PRN, nausea, vomiting Give if patient is able to tolerate oral medication.
<input checked="" type="checkbox"/> ondansetron (ZOFTRAN) 4 mg/2 mL injection	4 mg, intravenous, every 8 hours PRN, nausea, vomiting 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 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 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 Give if ondansetron (ZOFTRAN) is ineffective and patient is UNable to tolerate oral medication.

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

ramelteon (ROZEREM) tablet 8 mg, oral, nightly PRN, sleep

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

zolpidem (AMBIEN) tablet 5 mg, oral, nightly PRN, sleep

ramelteon (ROZEREM) tablet 8 mg, oral, nightly PRN, sleep

VTE

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

VTE/DVT Risk Definitions

URL:

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

[Anticoagulation Guide for COVID patients](#)

URL:

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

Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis with Risk Stratification (Single Response) (Selection Required)

Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)

Moderate risk of VTE Routine, Once

<input type="checkbox"/>	Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/>	Place sequential compression device (Single Response)	
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/>	Moderate risk of VTE	Routine, Once
<input type="checkbox"/>	Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/>	Place sequential compression device (Single Response)	
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/>	High risk of VTE	Routine, Once
<input type="checkbox"/>	Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/>	Place sequential compression device (Single Response)	
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/>	High risk of VTE	Routine, Once
<input type="checkbox"/>	Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/>	Place sequential compression device (Single Response)	
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	LOW Risk of DVT (Selection Required)	
	Low Risk Definition Age less than 60 years and NO other VTE risk factors	
<input type="checkbox"/>	Low Risk (Single Response) (Selection Required)	
<input type="checkbox"/>	Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation
<input type="checkbox"/>	MODERATE Risk of DVT - Surgical (Selection Required)	

Moderate Risk Definition

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

One or more of the following medical conditions:

CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome

Age 60 and above

Central line

History of DVT or family history of VTE

Anticipated length of stay GREATER than 48 hours

Less than fully and independently ambulatory

Estrogen therapy

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Moderate Risk (Selection Required)

Moderate risk of VTE Routine, Once

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

Contraindications exist for pharmacologic prophylaxis BUT order Sequential compression device **"And" Linked Panel**

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following contraindication(s):

Place/Maintain sequential compression device continuous Routine, Continuous

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

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following contraindication(s):

Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):

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

enoxaparin (LOVENOX) syringe 40 mg, subcutaneous, daily at 0600, Starting S+1
Indication(s): VTE Prophylaxis

patients with CrCL LESS than 30 mL/min 30 mg, subcutaneous, daily at 0600, Starting S+1
For Patients with CrCL LESS than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min 30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min 40 mg, subcutaneous, 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
Indication(s): VTE Prophylaxis

fondaparinux (ARIXTRA) injection 2.5 mg, subcutaneous, daily, Starting S+1
If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT):

heparin (porcine) injection 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM

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
Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.

<input type="checkbox"/>	HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM For patients with weight GREATER than 100 kg.
<input type="checkbox"/>	warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/>	Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)		
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> MODERATE Risk of DVT - Non-Surgical (Selection Required)		
Moderate Risk Definition		
Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated.		
One or more of the following medical conditions:		
CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome		
Age 60 and above		
Central line		
History of DVT or family history of VTE		
Anticipated length of stay GREATER than 48 hours		
Less than fully and independently ambulatory		
Estrogen therapy		
Moderate or major surgery (not for cancer)		
Major surgery within 3 months of admission		
<input type="checkbox"/> Moderate Risk (Selection Required)		
<input type="checkbox"/>	Moderate risk of VTE	Routine, Once
<input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)		
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis - Order Sequential compression device	"And" Linked Panel
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis	"And" Linked Panel
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)		
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S Indication(s): VTE Prophylaxis
<input type="checkbox"/>	patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 1700, Starting S For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/>	patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis

<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Selection Required)	
High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Enoxaparin for VTE Prophylaxis (Single Response)	
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Every 12 Hours	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	40 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Every 12 Hours	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	40 mg, subcutaneous, every 12 hours Indication(s):

<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, Starting S+1 For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Non-Surgical (Selection Required)	
High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 1700, Starting S For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis

<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
[] Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Hip/Knee) (Selection Required)	
High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE	
[] High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> aspirin chewable tablet	162 mg, oral, daily, Starting S+1
<input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet	162 mg, oral, daily, Starting S+1
<input type="checkbox"/> Apixaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> apixaban (ELIQUIS) tablet	2.5 mg, oral, 2 times daily, Starting S+1 Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor apixaban (ELIQUIS) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 Indication(s): VTE Prophylaxis

<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 140 kg or GREATER and CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM For patients with weight GREATER than 100 kg.
<input type="checkbox"/> Rivaroxaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission	10 mg, oral, daily at 0600 (TIME CRITICAL) Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor rivaroxaban (XARELTO) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous

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

VTE/DVT Risk Definitions

URL:

"\\appt1\epicapprod\Restricted\OrderSets\VTEDVTRISK DEFINITIONS.pdf"

[Anticoagulation Guide for COVID patients](#)

URL:

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

<input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis with Risk Stratification (Single Response) (Selection Required)	
<input type="checkbox"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/> Moderate risk of VTE	Routine, Once
<input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:

<input type="checkbox"/> Place sequential compression device (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/> Moderate risk of VTE	Routine, Once
<input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/> Place sequential compression device (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/> Place sequential compression device (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis	Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
<input type="checkbox"/> Place sequential compression device (Single Response)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> LOW Risk of DVT (Selection Required)	
Low Risk Definition Age less than 60 years and NO other VTE risk factors	
<input type="checkbox"/> Low Risk (Single Response) (Selection Required)	
<input type="checkbox"/> Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation
<input type="checkbox"/> MODERATE Risk of DVT - Surgical (Selection Required)	

Moderate Risk Definition

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

One or more of the following medical conditions:

CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome

Age 60 and above

Central line

History of DVT or family history of VTE

Anticipated length of stay GREATER than 48 hours

Less than fully and independently ambulatory

Estrogen therapy

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Moderate Risk (Selection Required)

Moderate risk of VTE Routine, Once

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

Contraindications exist for pharmacologic prophylaxis BUT order Sequential compression device **"And" Linked Panel**

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following contraindication(s):

Place/Maintain sequential compression device continuous Routine, Continuous

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

Contraindications exist for pharmacologic prophylaxis Routine, Once
No pharmacologic VTE prophylaxis due to the following contraindication(s):

Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):

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

enoxaparin (LOVENOX) syringe 40 mg, subcutaneous, daily at 0600, Starting S+1
Indication(s): VTE Prophylaxis

patients with CrCL LESS than 30 mL/min 30 mg, subcutaneous, daily at 0600, Starting S+1
For Patients with CrCL LESS than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min 30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
Indication(s): VTE Prophylaxis

patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min 40 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min
Indication(s): VTE Prophylaxis

fondaparinux (ARIXTRA) injection 2.5 mg, subcutaneous, daily, Starting S+1
If the patient does not have a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication. Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min. This patient has a history of or suspected case of Heparin-Induced Thrombocytopenia (HIT):

heparin (porcine) injection 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM

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
Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.

HEParin (porcine) injection - For Patients with weight GREATER than 100 kg 7,500 Units, subcutaneous, every 8 hours, Starting S+1
For patients with weight GREATER than 100 kg.

<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> MODERATE Risk of DVT - Non-Surgical (Selection Required)	
Moderate Risk Definition Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. One or more of the following medical conditions: CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome Age 60 and above Central line History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hours Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission	
<input type="checkbox"/> Moderate Risk (Selection Required)	
<input type="checkbox"/> Moderate risk of VTE	Routine, Once
<input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis - Order Sequential compression device	"And" Linked Panel
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis	"And" Linked Panel
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 1700, Starting S+1 For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis

<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours Recommended for patients with high risk of bleeding, e.g. weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Selection Required) Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Enoxaparin for VTE Prophylaxis (Single Response)	
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Every 12 Hours	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	40 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Every 12 Hours	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	40 mg, subcutaneous, every 12 hours Indication(s):
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.

<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, Starting S+1 For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> HIGH Risk of DVT - Non-Surgical (Selection Required)	
Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily, Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily, Starting S+1 For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, every 12 hours at 0900, 2100, Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, every 12 hours at 0900, 2100 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Hip/Knee) (Selection Required)	
Address both pharmacologic and mechanical prophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> aspirin chewable tablet	162 mg, oral, daily, Starting S+1
<input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet	162 mg, oral, daily, Starting S+1

<input type="checkbox"/> Apixaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> apixaban (ELIQUIS) tablet	2.5 mg, oral, 2 times daily, Starting S+1 Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor apixaban (ELIQUIS) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 140 kg or GREATER and CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM For patients with weight GREATER than 100 kg.
<input type="checkbox"/> Rivaroxaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission	10 mg, oral, daily at 0600 (TIME CRITICAL) Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor rivaroxaban (XARELTO) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:

DVT Risk and Prophylaxis Tool (Single Response)

VTE/DVT Risk Definitions

URL:

"\\appt1\epicapprod\Restricted\OrderSets\VTEDVTRISK DEFINITIONS.pdf"

URL:

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

- Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis with Risk Stratification (Single Response) (Selection Required)

- () Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)
- Moderate risk of VTE Routine, Once
- Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis Routine, Once
No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication.
Therapy for the following:
- Place sequential compression device (Single Response)
- () Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):
- () Place/Maintain sequential compression device continuous Routine, Continuous
- () Moderate Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)
- Moderate risk of VTE Routine, Once
- Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis Routine, Once
No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication.
Therapy for the following:
- Place sequential compression device (Single Response)
- () Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):
- () Place/Maintain sequential compression device continuous Routine, Continuous
- () High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)
- High risk of VTE Routine, Once
- Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis Routine, Once
No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication.
Therapy for the following:
- Place sequential compression device (Single Response)
- () Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):
- () Place/Maintain sequential compression device continuous Routine, Continuous
- () High Risk - Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis (Selection Required)
- High risk of VTE Routine, Once
- Patient currently has an active order for therapeutic anticoagulant or VTE prophylaxis Routine, Once
No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication.
Therapy for the following:
- Place sequential compression device (Single Response)
- () Contraindications exist for mechanical prophylaxis Routine, Once
No mechanical VTE prophylaxis due to the following contraindication(s):
- () Place/Maintain sequential compression device continuous Routine, Continuous
- () LOW Risk of DVT (Selection Required)
- Low Risk Definition
Age less than 60 years and NO other VTE risk factors

Low Risk (Single Response) (Selection Required)

<input type="checkbox"/> Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourage early ambulation
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MODERATE Risk of DVT - Surgical (Selection Required)

Moderate Risk Definition

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

One or more of the following medical conditions:

CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome

Age 60 and above

Central line

History of DVT or family history of VTE

Anticipated length of stay GREATER than 48 hours

Less than fully and independently ambulatory

Estrogen therapy

Moderate or major surgery (not for cancer)

Major surgery within 3 months of admission

Moderate Risk (Selection Required)

<input type="checkbox"/> Moderate risk of VTE	Routine, Once
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Moderate Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required)

<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis BUT order Sequential compression device	"And" Linked Panel
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<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
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<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
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<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis	"And" Linked Panel
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<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
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<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
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enoxaparin (LOVENOX) injection (Single Response) (Selection Required)

<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1 Indication(s): VTE Prophylaxis
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<input type="checkbox"/> patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
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<input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
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<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
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<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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):
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<input type="checkbox"/> heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
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<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> MODERATE Risk of DVT - Non-Surgical (Selection Required)	
Moderate Risk Definition Pharmacologic prophylaxis must be addressed. Mechanical prophylaxis is optional unless pharmacologic is contraindicated. One or more of the following medical conditions: CHF, MI, lung disease, pneumonia, active inflammation, dehydration, varicose veins, cancer, sepsis, obesity, previous stroke, rheumatologic disease, sickle cell disease, leg swelling, ulcers, venous stasis and nephrotic syndrome Age 60 and above Central line History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hours Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission	
<input type="checkbox"/> Moderate Risk (Selection Required)	
<input type="checkbox"/> Moderate risk of VTE	Routine, Once
<input type="checkbox"/> Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis - Order Sequential compression device	"And" Linked Panel
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis AND mechanical prophylaxis	"And" Linked Panel
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 1700, Starting S For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis

<input type="checkbox"/> patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Selection Required)	
High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Enoxaparin for VTE Prophylaxis (Single Response)	
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 30 mg Every 12 Hours	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	30 mg, subcutaneous, daily at 1700 Indication(s):
<input type="checkbox"/> enoxaparin (LOVENOX) 40 mg Daily at 1700	
<input type="checkbox"/> enoxaparin (LOVENOX) injection	40 mg, subcutaneous, daily at 1700 Indication(s):

<input type="checkbox"/>	enoxaparin (LOVENOX) 40 mg Every 12 Hours	
<input type="checkbox"/>	enoxaparin (LOVENOX) injection	40 mg, subcutaneous, every 12 hours Indication(s):
<input type="checkbox"/>	fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/>	HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, Starting S+1 For patients with weight GREATER than 100 kg.
<input type="checkbox"/>	warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/>	Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/>	Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/>	Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/>	HIGH Risk of DVT - Non-Surgical (Selection Required)	
	High Risk Definition	
	Both pharmacologic AND mechanical prophylaxis must be addressed.	
	One or more of the following medical conditions:	
	Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders)	
	Severe fracture of hip, pelvis or leg	
	Acute spinal cord injury with paresis	
	Multiple major traumas	
	Abdominal or pelvic surgery for CANCER	
	Acute ischemic stroke	
	History of PE	
<input type="checkbox"/>	High Risk (Selection Required)	
<input type="checkbox"/>	High risk of VTE	Routine, Once
<input type="checkbox"/>	High Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/>	Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/>	enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/>	enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S Indication(s): VTE Prophylaxis
<input type="checkbox"/>	patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 1700, Starting S For Patients with CrCL LESS than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/>	patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily, Starting S For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/>	patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis

<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours For patients with weight GREATER than 100 kg.
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous
<input type="checkbox"/> HIGH Risk of DVT - Surgical (Hip/Knee) (Selection Required)	
High Risk Definition Both pharmacologic AND mechanical prophylaxis must be addressed. One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin variant mutations, anticardiolipin antibody syndrome; antithrombin, protein C or protein S deficiency; hyperhomocysteinemia; myeloproliferative disorders) Severe fracture of hip, pelvis or leg Acute spinal cord injury with paresis Multiple major traumas Abdominal or pelvic surgery for CANCER Acute ischemic stroke History of PE	
<input type="checkbox"/> High Risk (Selection Required)	
<input type="checkbox"/> High risk of VTE	Routine, Once
<input type="checkbox"/> High Risk Pharmacological Prophylaxis - Hip or Knee (Arthroplasty) Surgical Patient (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> aspirin chewable tablet	162 mg, oral, daily, Starting S+1
<input type="checkbox"/> aspirin (ECOTRIN) enteric coated tablet	162 mg, oral, daily, Starting S+1
<input type="checkbox"/> Apixaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> apixaban (ELIQUIS) tablet	2.5 mg, oral, 2 times daily, Starting S+1 Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor apixaban (ELIQUIS) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) injection (Single Response) (Selection Required)	
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1 Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 Indication(s): VTE Prophylaxis

<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min. Indication(s): VTE Prophylaxis
<input type="checkbox"/> enoxaparin (LOVENOX) syringe - For Patients weight between 140 kg or GREATER and CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min Indication(s): VTE Prophylaxis
<input type="checkbox"/> fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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
<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 Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
<input type="checkbox"/> HEParin (porcine) injection - For Patients with weight GREATER than 100 kg	7,500 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM For patients with weight GREATER than 100 kg.
<input type="checkbox"/> Rivaroxaban and Pharmacy Consult (Selection Required)	
<input type="checkbox"/> rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this admission	10 mg, oral, daily at 0600 (TIME CRITICAL) Indications: VTE prophylaxis
<input type="checkbox"/> Pharmacy consult to monitor rivaroxaban (XARELTO) therapy	STAT, Until discontinued, Starting S Indications: VTE prophylaxis
<input type="checkbox"/> warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
<input type="checkbox"/> Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
<input type="checkbox"/> Mechanical Prophylaxis (Single Response) (Selection Required)	
<input type="checkbox"/> Contraindications exist for mechanical prophylaxis	Routine, Once No mechanical VTE prophylaxis due to the following contraindication(s):
<input type="checkbox"/> Place/Maintain sequential compression device continuous	Routine, Continuous

Labs

General

<input type="checkbox"/> CBC and differential	Once
<input type="checkbox"/> Hematocrit	Once, Starting S For 1 Occurrences
<input type="checkbox"/> Hemoglobin	Once
<input type="checkbox"/> Platelet count	Once
<input type="checkbox"/> Partial thromboplastin time	Once
<input type="checkbox"/> Prothrombin time with INR	Once
<input type="checkbox"/> Comprehensive metabolic panel	Once
<input type="checkbox"/> Basic metabolic panel	Once
<input type="checkbox"/> Hepatic function panel	Once
<input type="checkbox"/> BUN	Once
<input type="checkbox"/> Creatinine	Once
<input type="checkbox"/> Calcium	Once
<input type="checkbox"/> Ionized calcium	Once
<input type="checkbox"/> Phosphorus	Once

<input type="checkbox"/>	Magnesium	Once
<input type="checkbox"/>	Amylase	Once
<input type="checkbox"/>	Lipase	Once
<input type="checkbox"/>	Bilirubin, direct	Once
<input type="checkbox"/>	Bilirubin, total	Once
<input type="checkbox"/>	GGT	Once
<input type="checkbox"/>	C-reactive protein	Once
<input type="checkbox"/>	FK506 Tacrolimus level, random	Once
<input type="checkbox"/>	CLO test	Once, Biopsy
<input type="checkbox"/>	Cyclosporine level, random	Once
<input type="checkbox"/>	Protein electrophoresis, serum	Once
<input type="checkbox"/>	CK isoenzymes	Once
<input type="checkbox"/>	Erythropoietin	Once
<input type="checkbox"/>	Sedimentation rate	Once
<input type="checkbox"/>	Lactate dehydrogenase, LDH	Once
<input type="checkbox"/>	Syphilis treponema screen with RPR confirmation (reverse algorithm)	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):

General - HMSJ

<input type="checkbox"/>	CBC and differential	Once
<input type="checkbox"/>	Hematocrit	Once, Starting S For 1 Occurrences
<input type="checkbox"/>	Hemoglobin	Once
<input type="checkbox"/>	Platelet count	Once
<input type="checkbox"/>	Comprehensive metabolic panel	Once
<input type="checkbox"/>	Basic metabolic panel	Once
<input type="checkbox"/>	BUN	Once
<input type="checkbox"/>	Creatinine	Once
<input type="checkbox"/>	Hepatic function panel	Once
<input type="checkbox"/>	Amylase	Once
<input type="checkbox"/>	Erythropoietin	Once
<input type="checkbox"/>	Sedimentation rate	Once
<input type="checkbox"/>	Lactate dehydrogenase, LDH	Once
<input type="checkbox"/>	Lipase	Once
<input type="checkbox"/>	Bilirubin, direct	Once
<input type="checkbox"/>	Bilirubin, total	Once
<input type="checkbox"/>	GGT	Once
<input type="checkbox"/>	Magnesium	Once
<input type="checkbox"/>	Calcium	Once
<input type="checkbox"/>	Urinalysis with microscopic	Once
<input type="checkbox"/>	C-reactive protein	Once
<input type="checkbox"/>	FK506 Tacrolimus level, random	Once
<input type="checkbox"/>	CLO test	Once, Biopsy
<input type="checkbox"/>	Cyclosporine level, random	Once
<input type="checkbox"/>	Protein electrophoresis, serum	Once
<input type="checkbox"/>	Phosphorus	Once
<input type="checkbox"/>	CK isoenzymes	Once
<input type="checkbox"/>	Prealbumin	Once
<input type="checkbox"/>	Prothrombin time with INR	Once
<input type="checkbox"/>	Partial thromboplastin time	Once
<input type="checkbox"/>	PTH-related peptide	Once
<input type="checkbox"/>	Rheumatoid factor	Once
<input type="checkbox"/>	Syphilis treponema screen with RPR confirmation (reverse algorithm)	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):

Metabolic

<input type="checkbox"/>	Hemoglobin A1c	Once
<input type="checkbox"/>	Lipid panel	Once

<input type="checkbox"/>	Testosterone	Once
<input type="checkbox"/>	TSH	Once
<input type="checkbox"/>	T3	Once
<input type="checkbox"/>	T4	Once
<input type="checkbox"/>	Ferritin	Once
<input type="checkbox"/>	Iron	Once
<input type="checkbox"/>	Total iron binding capacity	Once
<input type="checkbox"/>	PSA	Once
		Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):
<input type="checkbox"/>	Transferrin	Once
<input type="checkbox"/>	Ceruloplasmin	Once
<input type="checkbox"/>	Cryo, globulin and fibrinogen	Once
<input type="checkbox"/>	Alpha-1-antitrypsin	Once
<input type="checkbox"/>	Alpha-1 antitrypsin phenotype	Once
<input type="checkbox"/>	Uric acid	Once
<input type="checkbox"/>	Vitamin A	Once
<input type="checkbox"/>	Vitamin B12	Once
<input type="checkbox"/>	Folate	Once
<input type="checkbox"/>	Vitamin D 25 hydroxy	Once
<input type="checkbox"/>	Vitamin E	Once
<input type="checkbox"/>	Zinc	Once
<input type="checkbox"/>	GGT	Once
<input type="checkbox"/>	Haptoglobin	Once
<input type="checkbox"/>	Troponin T	Once
<input type="checkbox"/>	CK total	Once
<input type="checkbox"/>	Carnitine, free and total	Once
<input type="checkbox"/>	B-type natriuretic peptide	Once
<input type="checkbox"/>	Ammonia	Once
<input type="checkbox"/>	Syphilis treponema screen with RPR confirmation (reverse algorithm)	Once
		Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):
<input type="checkbox"/>	Urinalysis screen and microscopy, with reflex to culture	Once
		Specimen Source: Urine
		Specimen Site:

Viral Workup - HMM

<input type="checkbox"/>	HIV Ag/Ab combination	Once
		Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):
<input type="checkbox"/>	Hepatitis A antibody, total	Once
<input type="checkbox"/>	Hepatitis A antibody, IgM	Once
<input type="checkbox"/>	Hepatitis B surface antibody	Once
<input type="checkbox"/>	Hepatitis B surface Ab, quantitative	Once
<input type="checkbox"/>	Hepatitis B surface antigen	Once
<input type="checkbox"/>	Hepatitis B core antibody, total	Once
<input type="checkbox"/>	Hepatitis B core antibody, IgM	Once
<input type="checkbox"/>	Hepatitis B e antibody	Once
<input type="checkbox"/>	Hepatitis B e antigen	Once
<input type="checkbox"/>	Hepatitis B virus (HBV), quantitative PCR	Once
<input type="checkbox"/>	Hepatitis C antibody	Once
<input type="checkbox"/>	Hepatitis C genotype	Once
<input type="checkbox"/>	Hepatitis C virus (HCV), quantitative PCR	Once
<input type="checkbox"/>	Hepatitis delta virus	Once
<input type="checkbox"/>	Hepatitis delta virus (HDV) Ab, IgM	Once

Viral Workup - HMSL/HMW

<input type="checkbox"/>	Cytomegalovirus antibody, IgG	Once
<input type="checkbox"/>	Cytomegalovirus antibody, IgM	Once

<input type="checkbox"/> Cytomegalovirus (CMV), PCR	Once Specimen Source: Plasma
<input type="checkbox"/> Cytomeg IgG/IgM	Once
<input type="checkbox"/> Epstein-Barr virus antibody test	Once
<input type="checkbox"/> Epstein Barr Virus (EBV) by PCR	Once Specimen Source: Plasma
<input type="checkbox"/> Rapid HIV 1 & 2	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):
<input type="checkbox"/> Hepatitis A antibody, total	Once
<input type="checkbox"/> Hepatitis A antibody, IgM	Once
<input type="checkbox"/> Hepatitis B core antibody, IgM	Once
<input type="checkbox"/> Hepatitis B core antibody, total	Once
<input type="checkbox"/> Hepatitis B e antibody	Once
<input type="checkbox"/> Hepatitis B e antigen	Once
<input type="checkbox"/> Hepatitis B surface antibody	Once
<input type="checkbox"/> Hepatitis B surface Ab, quantitative	Once
<input type="checkbox"/> Hepatitis B surface antigen	Once
<input type="checkbox"/> Hepatitis B virus (HBV), quantitative PCR	Once
<input type="checkbox"/> Hepatitis C antibody	Once
<input type="checkbox"/> Hepatitis C genotype	Once
<input type="checkbox"/> Hepatitis C virus (HCV), quantitative PCR	Once
<input type="checkbox"/> Hepatitis delta virus	Once
<input type="checkbox"/> Hepatitis delta virus (HDV) Ab, IgM	Once
<input type="checkbox"/> Hepatitis E virus Ab, IgG by ELISA	Once
<input type="checkbox"/> Hepatitis E virus Ab, IgM by ELISA	Once
<input type="checkbox"/> Herpes simplex virus, PCR	Once Specimen Source: Plasma

Viral Workup - HMCL, HMTW, HMSJ, HMWB

<input type="checkbox"/> Cytomegalovirus antibody, IgG	Once
<input type="checkbox"/> Cytomegalovirus antibody, IgM	Once
<input type="checkbox"/> Cytomegalovirus (CMV), PCR	Once Specimen Source: Plasma
<input type="checkbox"/> Cytomeg IgG/IgM	Once
<input type="checkbox"/> Epstein-Barr virus antibody test	Once
<input type="checkbox"/> Epstein Barr Virus (EBV) by PCR	Once Specimen Source: Plasma
<input type="checkbox"/> HIV 1, 2 antibody	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.):
<input type="checkbox"/> Hepatitis A antibody, total	Once
<input type="checkbox"/> Hepatitis A antibody, IgM	Once
<input type="checkbox"/> Hepatitis B core antibody, IgM	Once
<input type="checkbox"/> Hepatitis B core antibody, total	Once
<input type="checkbox"/> Hepatitis B e antibody	Once
<input type="checkbox"/> Hepatitis B e antigen	Once
<input type="checkbox"/> Hepatitis B surface antibody	Once
<input type="checkbox"/> Hepatitis B surface Ab, quantitative	Once
<input type="checkbox"/> Hepatitis B surface antigen	Once
<input type="checkbox"/> Hepatitis B virus (HBV), quantitative PCR	Once
<input type="checkbox"/> Hepatitis C antibody	Once
<input type="checkbox"/> Hepatitis C genotype	Once
<input type="checkbox"/> Hepatitis C virus (HCV), quantitative PCR	Once
<input type="checkbox"/> Hepatitis delta virus	Once
<input type="checkbox"/> Hepatitis delta virus (HDV) Ab, IgM	Once
<input type="checkbox"/> Hepatitis E virus Ab, IgG by ELISA	Once
<input type="checkbox"/> Hepatitis E virus Ab, IgM by ELISA	Once
<input type="checkbox"/> Herpes simplex virus, PCR	Once Specimen Source: Plasma

Autoimmune Workup

<input type="checkbox"/>	Anti smooth muscle Ab screen	Once
<input type="checkbox"/>	Gliadin peptide Abs, IgA and IgG	Once
<input type="checkbox"/>	Tissue transglutaminase, IgA	Once
<input type="checkbox"/>	Tissue transglutaminase, IgG	Once
<input type="checkbox"/>	Liver-kidney microsome Ab, IgG	Once
<input type="checkbox"/>	ANA	Once
<input type="checkbox"/>	Anti mitochondria screen	Once
<input type="checkbox"/>	Immunoglobulin G	Once
<input type="checkbox"/>	Immunoglobulin A	Once
<input type="checkbox"/>	Immunoglobulin M	Once
<input type="checkbox"/>	Immunoglobulin E	Once

Cancer Workup

<input type="checkbox"/>	Alpha fetoprotein	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.)
<input type="checkbox"/>	CEA	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.)
<input type="checkbox"/>	Cancer antigen 19-9	Once
<input type="checkbox"/>	CA 125	Once Release to patient (Note: If manual release option is selected, result will auto release 10 days from finalization.)
<input type="checkbox"/>	Chromogranin A	Once
<input type="checkbox"/>	Gastrin	Once

Copper Studies

<input type="checkbox"/>	Ceruloplasmin	Once
<input type="checkbox"/>	Copper, serum	Once
<input type="checkbox"/>	Copper, urine	Once

Stool Studies

<input type="checkbox"/>	Occult blood, stool	Conditional Frequency For 3 Occurrences, Stool When specimen available
<input type="checkbox"/>	Stool culture	Once For 1 Occurrences, Stool
<input type="checkbox"/>	Ova & Parasites-Concentrated Examination	Once, Stool
<input type="checkbox"/>	Fecal lactoferrin	Once, Stool
<input type="checkbox"/>	Giardia antigen	Once, Stool
<input type="checkbox"/>	Potassium, stool	Once, Starting S For 1 Occurrences, Stool
<input type="checkbox"/>	Sodium, stool	Once, Starting S For 1 Occurrences, Stool
<input type="checkbox"/>	Fecal fat, qualitative	Once, Stool
<input type="checkbox"/>	Cryptosporidium antigen, stool	Once, Stool
<input type="checkbox"/>	Porphyryn, total	Once
<input type="checkbox"/>	C difficile toxin / Gastrointestinal panels	
<input type="checkbox"/>	Enteric pathogen panels	
<input type="checkbox"/>	Enteric bacterial panel - Campylobacter spp., Enterotoxigenic E. coli (ETEC), Shigella spp./EIEC, Shiga-toxin producing E. coli (STEC), Salmonella spp., Shigella dysenteriae, P. shigelloides, Vibrio, Y. enterocolitica	Once, Stool
<input type="checkbox"/>	Enteric parasitic panel - G. lamblia, Cryptosporidium, E. histolytica	Once, Stool
<input type="checkbox"/>	Enteric viral panel - Norovirus GI & GII, Rotavirus A, Adenovirus F40/41, Sapovirus (genogroups I, II, IV, V), Human Astrovirus (hAstro)	Once, Stool

Patient has been an inpatient for greater than 3 days. C.difficile testing is appropriate for new onset diarrhea. Gastrointestinal Panel testing is not appropriate. If you have questions, please call the Microbiology Laboratory at 713-441-0330.

C difficile toxin Once, Stool
Reason to order:
Risk factors:

Patient has received a laxative, enema or medication with laxative effect. C. difficile or Gastrointestinal panel testing is not appropriate until laxative medication has been discontinued for 48 hours. If you have questions, please call the Microbiology Laboratory at 713-441-0330. (Selection Required)

@LAXPRINTGROUP@
@ENEMAPRINTGROUP@

Fecal lactoferrin Once, Stool

Fecal calprotectin Once, Stool

C difficile toxin (Single Response)

C difficile toxin Once, Stool
Reason to order:
Risk factors:

Gastrointestinal panel Once, Stool

Patient had a previous positive C. difficile / Gastrointestinal panel in the prior 14 days. Repeat C. difficile or Gastrointestinal panel testing is not appropriate. If you have questions, please call the Microbiology Laboratory at 713-441-0330.

@LASTLAB(CDIFFTOX,GASTROPANEL)@

Fecal lactoferrin Once, Stool

Fecal calprotectin Once, Stool

Enteric Precautions

Enteric isolation status Details

Misc Referral Lab Test

Hemochromatosis (HFE) 3 mutations Once

Hepatitis B virus DNA, Qualitative Once
Hepatitis B virus DNA, Qualitative

DCP (PIVKA II) Once
DCP (PIVKA II)

AFP-L3% Once
AFP-L3%

Microbiology

Blood culture x 2 **"And" Linked Panel**

Blood Culture (Aerobic & Anaerobic) Once, Blood
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, please call the lab for assistance; an IV line should NEVER be used.

Blood Culture (Aerobic & Anaerobic) Once, Blood
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, please call the lab for assistance; an IV line should NEVER be used.

Cardiology

Cardiology

Echocardiogram with agitated saline Routine, 1 time imaging, Starting S at 1:00 AM

Diagnostic Imaging

CT

CT Abdomen WWO Contrast (Omnipaque) **"And" Linked Panel**

For those with iodine allergies, please order the panel with Read-i-Cat (barium sulfate).

CT Abdomen W Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1
Liver protocol

iohexol (OMNIPAQUE) 300 mg iodine/mL
oral solution 30 mL, oral, once

CT Chest Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1

MRI/MRA

MRI Abdomen W Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1
Gadolinium contrast

MRI Abdomen W Wo Contrast Routine, 1 time imaging, Starting S at 1:00 AM For 1
EOVIST contrast

MRI Cholangiogram Routine, 1 time imaging, Starting S at 1:00 AM For 1

MRI Bone Survey Routine, 1 time imaging, Starting S at 1:00 AM For 1

X-Ray

CHEST 2 VW Routine, 1 time imaging, Starting S at 1:00 AM For 1

Upper GI and Small Bowel Routine, 1 time imaging, Starting S at 1:00 AM For 1

US

US Abdomen Complete Routine, 1 time imaging, Starting S at 1:00 AM For 1

US Abdominal Limited Routine, 1 time imaging, Starting S at 1:00 AM For 1

US Hepatic Routine, 1 time imaging, Starting S at 1:00 AM For 1

US Abdominal Doppler Routine, 1 time imaging, Starting S at 1:00 AM For 1

Other Diagnostic Studies

Other Diagnostic Studies

NM Hepatobiliary Routine, 1 time imaging, Starting S at 1:00 AM For 1

NM Gastric Emptying Routine, 1 time imaging, Starting S at 1:00 AM For 1
4 hours

IR Consult To Interventional Radiology Routine, 1 time imaging, Starting S at 1:00 AM For 1
Transjugular liver biopsy with portal pressure measurements.

Image Guidance Biopsy Routine, 1 time imaging, Starting S at 1:00 AM For 1

Consults

Physician Consults

Consult Hepatology Reason for Consult?
Patient/Clinical information communicated?
Patient/clinical information communicated?

Ancillary Consults

Consult to case management Consult Reason:

Consult to social work Reason for Consult:

PT eval and treat Reasons for referral to Physical Therapy (mark all applicable):
Are there any restrictions for positioning or mobility?
Please provide safe ranges for HR, BP, O2 saturation(if
values are very abnormal):
Weight Bearing Status:

Consult PT wound care Special Instructions:
Location of Wound?

[] OT eval and treat	Reason for referral to Occupational Therapy (mark all that apply): Are there any restrictions for positioning or mobility? Please provide safe ranges for HR, BP, O2 saturation(if values are very abnormal): Weight Bearing Status:
[] Consult to Nutrition	Reason For Consult? Purpose/Topic:
[] Consult to Spiritual Care	Reason for consult?
[] Consult to Speech Language Pathology	Routine, Once Reason for consult:
[] Consult to Wound Ostomy Care Nurse	Reason for consult: Reason for consult: Reason for consult: Reason for consult: Consult for NPWT: Reason for consult: Reason for consult:
[] Consult transplant social work	Reason for Consult? Organ Transplant: