Present on Admission (Single Response) (Se	election Required)
() COVID-19 virus detected	Details
() Suspected COVID-19 Virus	Details
Admission (Single Response) Patient has active status order on file.	
() Admit to inpatient	Diagnosis: Admitting Physician: Level of Care: Patient Condition: Bed request comments: Certification: I certify that based on my best clinical judgment and the patient's condition as documented in the HP and progress notes, I expect that the patient will need hospital services for two or more midnights.
Code Status	
[] Full code	Code Status decision reached by:
[] DNR (Selection Required)	
[] DNR (Do Not Resuscitate)	Does patient have decision-making capacity?
[] Consult to Palliative Care Service	Priority: Reason for Consult? Order?
	Name of referring provider:
[] Consult to Social Work	Name of referring provider:
[] Consult to Social Work [] Modified Code	Name of referring provider: Enter call back number:

COVID-19 ISOLATION NOT REQ

	ACUTE CARE PATIENT WITH NO AEROSOL GENERATING PROCEDURES	PATIENT WITH INTERMITTENT AEROSOL GENERATING TREATMENT/PROCEDURES	CRITICAL CARE PATIENT WITH CONTINUOUS AEROSOL GENERATING TREATMENT/PROCEDURES
Precautions	Standard + Droplet + Contact + Eye Protection	Standard + Modified Droplet + Contact + Eye Protection	Standard + Airborne + Contact + Eye Protection

[] Acute care patient with no aerosol generating procedures	
Droplet isolation status	Include eye protection
[] Contact isolation status	Include eye protection
[] Patient with intermittent aerosol generating	
treatment/procedures	
[] Modified droplet isolation status	Include eye protection
[] Contact isolation status	Include eye protection
[] Critical care patient with continuous aerosol gen	erating
treatment/procedures	
[] Airborne isolation status	Include eye protection

[] Contact isolation status

Precautions

[] Aspiration precautions	Details
[] Fall precautions	Increased observation level needed:
[] Latex precautions	Details
[] Seizure precautions	Increased observation level needed:

Nursing

Vital Signs (Selection Required) Vital signs with link to algorithm of Stepwise management of Hypoxemia

[X] Vital signs - T/P/R/BP	Routine, Per unit protocol For Until specified
[X] Pulse oximetry continuous	Routine, Continuous For Until specified Current FIO2 or Room Air:
Nursing Care	
[X] Strict intake and output for a target of 24 hour N EVEN balance	ET Routine, Every hour
[X] Limit repeated entry to room	Routine, Until discontinued, Starting S For Until specified Batch all care and work with pharmacy and providers to limit repeated entry to patient care room.
[X] Oral care for intubated patients	Routine, Every 4 hours For intubated patients
[X] Oral care for non intubated patients	Routine, Every shift For non intubated patients
[] Hemodynamic Monitoring	Routine, Continuous Measure:
[] Measure central venous pressure	Routine, Every 4 hours
[] Telemetry	"And" Linked Panel
[] Telemetry monitoring [] Telemetry Additional Setup Information	Routine, Continuous Order: Place in Centralized Telemetry Monitor: EKG Monitoring Only (Telemetry Box) Reason for telemetry: Can be off of Telemetry for tests and baths? Yes 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
[] Foley-NOT Recommended if patient able to void	
[] Insert Foley catheter	Routine, Once Type: Temperature Sensing Size: Urinometer needed:
[] Foley Catheter Care	Routine, Until discontinued, Starting S Orders: Maintain
[] Neurological assessment	Routine, Every shift Assessment to Perform:
[] Peripheral vascular assessment	Routine, Every 6 hours

Head of bed: 30 degrees Routine, Daily
····, _ ···;
Routine, Until discontinued, Starting S
Routine, Until discontinued, Starting S
Bathroom Privileges: with bathroom privileges
Routine, Until discontinued, Starting S
Specify: Up with assistance
Routine, Until discontinued, Starting S Specify: Activity as tolerated
Routine, Until discontinued, Starting S
Routine, Until discontinued, Starting S
, , , ,
Routine, Until discontinued, Starting S
Temperature greater than:
Temperature less than:
Systolic BP greater than:
Systolic BP less than:
Diastolic BP greater than:
Diastolic BP less than:
MAP less than: 65
Heart rate greater than (BPM): 120
Heart rate less than (BPM): 60
Respiratory rate greater than:
Respiratory rate less than: SpO2 less than: 92
Routine, Until discontinued, Starting S For Until specified, For
critical values.
Diet effective now, Starting S
NPO: Pre-Operative fasting options:
Diet effective now, Starting S
NPO: Except meds
Pre-Operative fasting options:
Diet effective now, Starting S
Diet(s):
Other Options:
Advance Diet as Tolerated?
Liquid Consistency:
Fluid Restriction:
Foods to Avoid:

() Tube Feeding

	Tube feeding - continuous	Continuous Tube Feeding Formula: Tube Feeding Formula: Tube Feeding Formula: Tube Feeding Formula: Tube Feeding Formula: Tube Feeding Schedule: Continuous Tube Feeding Route: Initial Tube Feed rate (mL/hr): Advance Rate by (mL/hr): Goal Tube Feed Rate (mL/hr): Dietitian to manage Tube Feed?
[]	XR Abdomen 1 Vw	Routine, 1 time imaging For 1

IV Fluids-IV Fluids for COVID-19 Should be Minimized

IV Fluids for COVID-19 Should be Minimized

Insert and Maintain IV / Central Line Access

[X] Insert and Maintain IV	"And" Linked Panel
[X] Insert peripheral IV	STAT, Once For 1 Occurrences
[X] Saline lock IV	Routine, Once For 1 Occurrences
[X] sodium chloride 0.9 % flush	10 mL, intravenous, PRN, line care
] Consult for Venous Access	Access:
	If GFR less than 45, has nephrology been consulted?
Bolus Fluids (Single Response) () sodium chloride 0.9 % bolus 500 mL	500 mL, intravenous, for 15 Minutes, once, For 1 Doses
) sodium chloride 0.9 % bolus 1000 mL	1,000 mL, intravenous, for 30 Minutes, once, For 1 Doses
) lactated ringer's bolus 500 mL	500 mL, intravenous, for 15 Minutes, once, For 1 Doses
) lactated ringers bolus 1000 mL	1,000 mL, intravenous, for 30 Minutes, once, For 1 Doses
() albumin human 5 % bottle	25 g, intravenous, for 15 Minutes, once, For 1 Doses Indication:

Medications

Pharmacy Consults

[X] Pharmacy consult to change IV medications to	STAT, Until discontinued, Starting S
concentrate fluids maximally	
[X] Pharmacy consult to manage dose adjustments for renal	STAT, Until discontinued, Starting S
function	Adjust dose for:

General COVID-19 Treatment

HM actively DISCOURAGES the combination of Azithromycin (+) Hydroxychloroquine as a treatment for COVID -19. The use of Hydroxychloroquine/Chloroquine for COVID-19 at HM should only be done in the context of a clinical trial. Contact Clinical Pharmacy with questions.

Screen patients for benefit of inclusion in HM COVID Investigational Protocols

%20CO URL: "ht URL: "https://f	parchives.com/houstonmethodist/documents/HM /ID%20algorithm.pdf" tps://covidtrials.houstonmethodist.org/" parchives.com/houstonmethodist/documents/COV 20Anticoagulation%20Guideline.pdf"
-----------------------------------------	-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

[] lopinavir-ritonavir (KALETRA) (Single Response)

() lopinavir-ritonavir (KALETRA) tablet	2 tablet, oral, 2 times daily Reason for Therapy: Viral Infection Documented Indication: Other Specify: COVID-19 New initiation of treatment for COVID-19 therapies are RESTRICTED to Infectious Diseases, Pulmonology and Critical Care Medicine providers. Are you an ID, Pulmonology or Critical Care provider?
() lopinavir-ritonavir (KALETRA) oral solution	5 mL, oral, 2 times daily Reason for Therapy: Viral Infection Documented Indication: Other Specify: COVID-19 New initiation of treatment for COVID-19 therapies are RESTRICTED to Infectious Diseases, Pulmonology and Critical Care Medicine providers. Are you an ID, Pulmonology or Critical Care provider?
Pharmacy consult for remdesivir Remdesivir via the FDA's EUA is subject to drug a	availability at HM
	URL: "file://\appt1.pdf"
	URL:
	"https://fparchives.com/houstonmethodist/documents/HM
	%20EUA%20RDV%20Criteria.pdf"
[] Pharmacy consult for remdesivir	Routine, Until discontinued, Starting S
	Physician contact number:
	Remdesivir prescribing is RESTRICTED to Infectious
	Diseases, Pulmonology and Critical Care Medicine providers.
	Are you an ID, Pulmonology or Critical Care provider?
Immunomodulatory Agents	
[] tocilizumab (ACTEMRA) IVPB	400 mg, intravenous, once, For 1 Doses RESTRICTED to infectious diseases, pulmonary, or critical
	care specialists. Are you a specialist or ordering on behalf of
	one?
[] anakinra (KINERET) subcutaneous syringe	100 mg, subcutaneous, every 8 hours, For 9 Doses
[] inFLIXimab (REMICADE) IVPB	5 mg/kg, intravenous, for 120 Minutes, once, For 1 Doses
Antibiotics	
[] azithromycin (ZITHROMAX) IV	intravenous, for 60 Minutes
	Reason for Therapy:
[] cefepime (MAXIPIME) IV	intravenous Reason for Therapy:
[] cefTRIAxone (ROCEPHIN) IV	intravenous, for 30 Minutes
[] cefTRIAxone (ROCEPHIN) IV	Reason for Therapy:
[] linezolid (ZYVOX) IV	intravenous, for 60 Minutes, every 12 hours
	Reason for Therapy:
[] piperacillin-tazobactam (ZOSYN) IV	intravenous
	Reason for Therapy:
[] meropenem (MERREM) IV	500 mg, intravenous, every 6 hours
	Reason for Therapy:
[] metronidazole (FLAGYL) IV	intravenous
	Reason for Therapy:
[] vancomycin (VANCOCIN) IV (Single Response)	
() vancomycin (VANCOCIN) IV - for	intravenous
PERIPHERAL LINE USE ONLY	Reason for Therapy:
() vancomycin (VANCOCIN) IV - for	intravenous
CENTRAL LINE USE ONLY	Reason for Therapy:
I Contraction of the second	

Scheduled Antihypertensives (Single Response)

) labetalol (NORMODYNE) tablet	200 mg, oral, 2 times daily at 0600, 1800 Hold Parameters: Hold Parameters Requested	
	Specify: Hold for heart rate less than 60/min or if s	ystolic
	blood pressure is less than 100 mmHg. Contact Physician:	
) labetalol (NORMODYNE)	intravenous, 2 times daily at 0600, 1800	
	HOLD parameters for this order: Hold Parameters HOLD for: Other Systolic BP	requested
	HOLD for: Other Heart Rate	
	Hold for Heart Rate LESS than (in bpm): 55	
	Contact Physician if: Hold for Systolic BP LESS than (in mmHg): 110	
) metoprolol tartrate (LOPRESSOR) tablet	100 mg, oral, 2 times daily at 0600, 1800	
	Hold Parameters: Hold Parameters Requested Specify: Hold for heart rate less than 60/min or if s	vstolic
	blood pressure is less than 100 mmHg.	,
) metoprolol (LOPRESSOR) injection	Contact Physician:	
) metoprolol (LOPRESSOR) injection	5 mg, intravenous Hold for heart rate less than 60/min or if systolic bl	bod
	pressure isless than 100 mmHg.	
	HOLD parameters for this order:	
) hydrALAZINE (APRESOLINE) injection	Contact Physician if: 10 mg, intravenous, every 6 hours	
) · · · · · · · · · · · · · · · · · · ·	If systolic blood pressure is greater than ***.	
	HOLD parameters for this order:	
	Contact Physician if:	
PRN Antihypertensives		
] labetalol (NORMODYNE, TRANDATE) injection -		
an alternative agent if heart rate is LESS than 55	3PM Systolic Blood Pressure GREATER than 160 mmH Administer at 2 mg/minute.	lg
	HOLD for: Other Heart Rate	
	Hold for Heart Rate LESS than (in bpm): 55	
	Contact Physician if: native 10 mg, intravenous, every 6 hours PRN, high bloo	-l
] hydrALAZINE (APRESOLINE) injection - Use alte		a pressure,
therapy if patient is tachycardic (GREATER than		
therapy if patient is tachycardic (GREATER than BPM)		lg
	00 Systolic Blood Pressure GREATER than 160 mmH	lg
BPM)	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE	lg
BPM)	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if:	lg
BPM)	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if:	lg
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ad	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel	lg
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous 	lg R than 100
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN C	lg R than 100
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous 	lg R than 100
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN C	lg R than 100
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	00 Systolic Blood Pressure GREATER than 160 mmH Hold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel illure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN C MEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate infusion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe	lg R than 100 DRDER ery hour to at TOF in
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN CMEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate inf usion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusion 	lg R than 100 DRDER ery hour to at TOF in on rate by 0
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN CMEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate infusion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusion mcg/kg/min. Monitor TOF every hour to achieve and maintair 	lg R than 100 DRDER ery hour to at TOF in on rate by (a 2 of 4 TO
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN C MEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate infusion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusio mcg/kg/min. Monitor TOF every hour to achieve and maintair Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, the same infusion rate, then repeat TOF in 4 hours. IF TOF L 	lg R than 100 DRDER ery hour to at TOF in on rate by 0 2 of 4 TO CONTINU ESS than
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN CMEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate inf usion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusion mcg/kg/min. Monitor TOF every hour to achieve and maintair Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, the same infusion rate, then repeat TOF in 4 hours. IF TOF L of 4, DECREASE infusion rate by 0.5 mcg/kg/min. Monitor TOF 	Ig R than 100 RDER ery hour to at TOF in on rate by 0 CONTINU ESS than OF every
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN CMEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate inf usion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusion mcg/kg/min. Monitor TOF every hour to achieve and maintair Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, the same infusion rate, then repeat TOF in 4 hours. IF TOF L of 4, DECREASE infusion rate by 0.5 mcg/kg/min. Monitor TOF hour to achieve and maintain 2 of 4 TOF. 	Ig R than 100 RDER ery hour to at TOF in on rate by 0 2 of 4 TOI CONTINU ESS than 2 OF every
BPM) Neuromuscular Blockage (Single Response) Dose based on Ideal body weight (IBW), unless ac) cisatracurium (NIMbex) Continuous Infusion Recommended for patients with renal or hepatic f	 O0 Systolic Blood Pressure GREATER than 160 mmHold for Heart Rate LESS than (in bpm): GREATE Contact Physician if: ual body weight LESS than ideal body weight. "Followed by" Linked Panel ilure. 1-10 mcg/kg/min, intravenous, continuous **PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN CMEDICATION DOSED BY IDEAL BODY WEIGHT** Initiate inf usion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min ev achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repe four hours. IF TOF GREATER than 2 of 4, INCREASE infusion mcg/kg/min. Monitor TOF every hour to achieve and maintair Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, the same infusion rate, then repeat TOF in 4 hours. IF TOF L of 4, DECREASE infusion rate by 0.5 mcg/kg/min. Monitor TOF 	Ig R than 100 RDER ery hour to at TOF in on rate by 0 2 of 4 TOI CONTINU ESS than 1 OF every

[] cisatracurium (NIMbex) injection	0.15 mg/kg, intravenous, once, For 1 Doses
[] cisatracurium (NIMbex) infusion	1-10 mcg/kg/min, intravenous, continuous
	**PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN ORDER
	MEDICATION DOSED BY IDEAL BODY WEIGHT**
	···· · · · · · · · · · · · · · · · · ·
	Initiate infusion at 1mcg/kg/min. Titrate by 0.5 mcg/kg/min every hour to
	achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repeat TOF in
	four hours. IF TOF GREATER than 2 of 4, INCREASE infusion rate by 0.5
	mcg/kg/min. Monitor TOF every hour to achieve and maintain 2 of 4 TOF.
	Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, CONTINUE
	the same infusion rate, then repeat TOF in 4 hours. IF TOF LESS than 2
	of 4, DECREASE infusion rate by 0.5 mcg/kg/min. Monitor TOF every
	hour to achieve and maintain 2 of 4 TOF. Once at 2 of 4 TOF, repeat
	TOF in four hours. Max dose 10mcg/kg/min.
() vecuronium (NORCURON) Continuous Infusion	"Followed by" Linked Panel
Use caution in patients with renal or hepatic dysfu	Inction
vecuronium (NORCURON) 1 mg/mL in	0.8-1.5 mcg/kg/min, intravenous, continuous
sodium chloride 0.9% 100 mL infusion	**PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN ORDER
	MEDICATION DOSED BY IDEAL BODY WEIGHT**
	Initiate infusion at 0.8mcg/kg/min. Titrate by 0.1 mcg/kg/min every hour to
	achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repeat TOF in
	four hours. IF TOF GREATER than 2/4, INCREASE infusion rate by 0.1
	mcg/kg/min. Monitor TOF every hour to achieve and maintain 2 of 4 TOF.
	Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, CONTINUE
	the same infusion rate, then repeat TOF in 4 hours. IF TOF LESS than 2
	of 4, DECREASE infusion rate by 0.1 mcg/kg/min. Monitor TOF every
	hour to achieve and maintain 2 of 4 TOF. Once at 2 of 4 TOF, repeat
	TOF in four hours. Max dose 1.5mcg/kg/min.
() vecuronium (NORCURON) IV Bolus and Continue	
Infusion	
Use caution in patients with renal or hepatic dysfu	Inction
[] vecuronium (NORCURON) injection	0.1 mg/kg, intravenous, once, For 1 Doses
[] vecuronium (NORCURON) 1 mg/mL in	0.8-1.5 mcg/kg/min, intravenous, continuous
sodium chloride 0.9% 100 mL infusion	**PROGRAM INFUSION PUMP WITH WEIGHT NOTED IN ORDER
	MEDICATION DOSED BY IDEAL BODY WEIGHT**
	Initiate infusion at 0.8mcg/kg/min. Titrate by 0.1 mcg/kg/min every hour to
	achieve 2 of 4 Train of Four (TOF). Once at 2 of 4 TOF, repeat TOF in
	four hours. IF TOF GREATER than 2/4, INCREASE infusion rate by 0.1
	mcg/kg/min. Monitor TOF every hour to achieve and maintain 2 of 4 TOF.
	Once at 2 of 4 TOF, repeat TOF in four hours. IF TOF 2 of 4, CONTINUE
	the same infusion rate, then repeat TOF in 4 hours. IF TOF LESS than 2
	of 4, DECREASE infusion rate by 0.1 mcg/kg/min. Monitor TOF every
	hour to achieve and maintain 2 of 4 TOF. Once at 2 of 4 TOF, repeat
	TOF in four hours. Max dose 1.5mcg/kg/min.
Vasoactive Drips	
DOPamine (INTROPIN) infusion	2-20 mcg/kg/min, intravenous, titrated
DOButamine (DOBUTREX) infusion	0.5-20 mcg/kg/min, intravenous, titrated
[] EPINEPHrine (ADRENALIN) in sodium chloride 0 250 mL infusion	.9 % 2-30 mcg/min, intravenous, titrated
	2-30 mcg/min introvonous titrated
	2-30 mcg/min, intravenous, titrated
[] phenylephrine (NEO-SYNEPHRINE) in sodium ch	nloride 5-150 mcg/min, intravenous, titrated
0.9 % 250 mL infusion	
[] vasopressin (PITRESSIN) 0.4 Units/mL in sodium	0.04 Units/min, intravenous, titrated
chloride 0.9 % 100 mL infusion	
[] milrinone infusion 200 mcg/mL (premixed)	0.125-0.75 mcg/kg/min, intravenous, titrated
[] nitroglycerin infusion	5-200 mcg/min, intravenous, titrated
[] nitroprusside (NIPRIDE) infusion	0.3-8 mcg/kg/min, intravenous, titrated

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[] niCARdipine (CARDENE) IV infusion	2.5-15 mg/hr, intravenous, titrated
[] esmolol (BREVIBLOC) infusion	50-200 mcg/kg/min, intravenous, titrated
Sedation	
] propofol (DIPRIVAN) or DEXMEDETomidine (PREcedex) infusion	
[] propofol (DIPRIVAN) infusion	0-50 mcg/kg/min, intravenous, continuous After initiation reassess RASS/BIS within 10 min. Titrate for Sedation. LESS than desired sedation effect: INCREASE rate by 5 mcg/kg/min. Reassess sedation within 10 minutes. DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. GREATER than desired sedation effect: DECREASE rate 5 mcg/kg/min and reassess sedation within 15 minutes. If patient requiring GREATER than: 50 mcg/kg/min, Contact MD to re evolute coddition therapy.
[] dexMEDEtomidine (PREcedex) infusion	 re-evaluate sedation therapy 0.1-1.5 mcg/kg/hr, intravenous, continuous Generally for mild to moderate sedation. Not for use in patients on neuromuscular blocking agents. NO LOADING DOSE. After initiation reassess RASS within 1 hour. Titrate for Sedation. LESS than desired sedation effect: INCREASE rate by 0.1 mcg/kg/hour. Reassess RASS within 1 hours. DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours GREATER than desired sedation effect: DECREASE rate by 0.1 mcg/kg/hour. If patient requiring GREATER than: 1.5 mcg/kg/hr, Contact MD to re-evaluate sedation therapy
] lorazepam (ATIVAN) or midazolam (VERSED) ir NOT HMW (Single Response)	
() lorazepam (ATIVAN) 60 mg/30 mL infusion	Loading Dose (optional): Not Ordered Nursing Bolus Dose: 0.5 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose IVOnce and increase rate by 0.25 milligram/hour then reassess sedation inone hour. If DESIRED sedation effect: Continue the same rate. Reassess sedationwithin 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation withi one hour. If patient requires GREA TER than 5 milligram/hour lorazepam contact MDto re-evalute sedation therapy. Indication(s): Sedation
() midazolam (VERSED) 60 mg/30 mL infusion	intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose IVOnce and increase rate by 0.25 milligram/hour then reassess sedation inone hour. If DESIRED sedation effect: Continue the same rate. Reassess sedationwithin 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation withi one hour. If patient requires GREA TER than 5 milligram/hour midazolan contact MDto re-evalute sedation therapy. Indication(s): Sedation

HMWB Only (Single Response)

()	LORAZepam (ATIVAN) 60 mg/30 mL infusion	Loading Dose (optional): Not Ordered Nursing Bolus Dose: 0.5 mg Continuous Dose: Not Ordered intravenous, continuous
	If LESS than desired sedation effect: administer ordered BOLUS dose IV	
		Once and increase rate by 0.25 milligram/hour then reassess sedation in
		one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation
		within 4 hours.
		If GREATER than desired sedation effect: Decrease rate by 0.25
		milligram/hour and reassess sedation within one hour.
		If patient requires GREA TER than 5 milligram/hour lorazepam, contact
		MD
		to re-evalute sedation therapy.
		Indication(s): Sedation
$\overline{()}$	MIDAZolam (VERSED) 30 mg/30 mL	intravenous, continuous
.,	infusion	If LESS than desired sedation effect: administer ordered BOLUS dose I'
		Once and increase rate by 0.25 milligram/hour then reassess sedation i
		one hour.
		If DESIRED sedation effect: Continue the same rate. Reassess sedation
		within 4 hours.
		If GREATER than desired sedation effect: Decrease rate by 0.25
		milligram/hour and reassess sedation within one hour.
		If patient requires GREA TER than 5 milligram/hour midazolam, contac
		MD
		to re-evalute sedation therapy. Indication(s): Sedation
	razepam (ATIVAN) or midazolam (VERSED)	
н	IVIVY UNIV (SINGLE RESDONSE)	
	MW Only (Single Response) LORAZepam (ATIVAN) 30 mg/30 mL	Loading Dose (optional): Not Ordered Nursing Bolus Dose: 0.5
	LORAZepam (ATIVAN) 30 mg/30 mL infusion	Loading Dose (optional): Not Ordered Nursing Bolus Dose: 0.5 mg Continuous Dose: Not Ordered
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour.
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedati
	LORAZepam (ATIVAN) 30 mg/30 mL	mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedati within 4 hours.
	LORAZepam (ATIVAN) 30 mg/30 mL	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedati within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25
	LORAZepam (ATIVAN) 30 mg/30 mL	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour.
	LORAZepam (ATIVAN) 30 mg/30 mL	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact
	LORAZepam (ATIVAN) 30 mg/30 mL	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD
	LORAZepam (ATIVAN) 30 mg/30 mL	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy.
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation is one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour.
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()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour.
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation is one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedati within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reasses sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour.
	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedati within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour midazolam, contact MD
()	LORAZepam (ATIVAN) 30 mg/30 mL infusion MIDAZolam in 0.9% NaCl (VERSED) 55	 mg Continuous Dose: Not Ordered intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If patient requires GREA TER than 5 milligram/hour lorazepam, contact MD to re-evalute sedation therapy. Indication(s): Sedation intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose I Once and increase rate by 0.25 milligram/hour then reassess sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reasses sedation i one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation i one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour. If GREATER than desired sedation effect: Decrease rate by 0.25 milligram/hour and reassess sedation within one hour.

infusion - HMSJ Only (Single Response)

() fentaNYL (SUBLIMAZE) 1500 mcg/30 mL infusion	intravenous, continuous **Due to fentaNYL 600 mcg/30 mL shortages, the new standard for all facilities will be fentaNYL 1500 mcg/30 mL. This concentration is 2.5 x more concentrated.**
	If LESS than desired sedation effect: administer ordered BOLUS dose IV Once and increase rate by 25 micrograms/hour then reassess sedation in one hour.
	If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 25 micrograms/hour and reassess sedation within one hour. If patient requires GREATE R than 200 micrograms/hour fentanyl, contac MD
	to re-evalute sedation therapy.
() hydromorPHONE (DILAUDID) 30 mg/30 mL in sodium chloride 0.9% infusion	intravenous, continuous If LESS than desired sedation effect: administer ordered BOLUS dose IV Once and increase rate by 0.5 milligrams/hour then reassess sedation in one hour. If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours.
	If GREATER than desired sedation effect: Decrease rate by 0.5 milligrams/hour and reassess sedation within one hour.
	If patient requires GREA TER than 2 milligrams/hour hydromorphone, contact
	MD to re-evalute sedation therapy.
[] fentanyl (SUBLIMAZE) or hydromorPHONE (DIL	AUDID)
infusion - NOT HMSJ (Single Response) () fentaNYL (SUBLIMAZE) 1500 mcg/30 mL	intravenous, continuous
infusion	**Due to fentaNYL 600 mcg/30 mL shortages, the new standard for all facilities will be fentaNYL 1500 mcg/30 mL. This concentration is 2.5 x more concentrated.**
	If LESS than desired sedation effect: administer ordered BOLUS dose IV Once and increase rate by 25 micrograms/hour then reassess sedation ir one hour.
	If DESIRED sedation effect: Continue the same rate. Reassess sedation within 4 hours. If GREATER than desired sedation effect: Decrease rate by 25
	micrograms/hour and reassess sedation within one hour. If patient requires GREATE R than 200 micrograms/hour fentanyl, contac MD
() hydromorPHONE (DILAUDID) 15 mg/30 mL	to re-evalute sedation therapy.
infusion	If LESS than desired sedation effect: administer ordered BOLUS dose IVOnce and increase rate by 0.5 milligrams/hour then reassess sedation inone hour. If DESIRED sedation effect: Continue the same rate. Reassess sedationwithin 4 hours. If GREATER than desired sedation effect: Decrease rate by 0.5 milligrams/hour and reassess sedation within one hour. If patient requires GREA TER than 2 milligrams/hour hydromorphone, contactMD to re-evalute sedation therapy.
Antitussives (Single Response)	
() guaiFENesin (MUCINEX) 12 hr tablet	1,200 mg, oral, every 12 hours PRN, cough
 () guai Livesii (MOCINEX) 12 in tablet () dextromethorphan-guaifenesin (ROBITUSSIN-D 10-100 mg/5 mL liquid 	
() benzonatate (TESSALON) capsule	200 mg, oral, every 8 hours PRN, cough
Antipyretics	
[] acetaminophen (TYLENOL) tablet	500 mg, oral, every 4 hours PRN, fever, Fever GREATER than 100.5 F
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[] acetaminophen (OFIRMEV) injection

1,000 mg, intravenous, for 15 Minutes, once, For 1 Doses IV acetaminophen (Ofirmev) is restricted to use only in OR, PACU, or ICU areas, and for patients that cannot tolerate oral, per tube, or rectal routes of administration. Do you attest that this restriction has been met?

Stress Ulcer Prophylaxis (Single Response)

Stress Ulcer Prophylaxis (Single Response)	
() famotidine (PEPCID) IV or ORAL	"Or" Linked Panel
[] famotidine (PEPCID) injection	20 mg, intravenous, every 12 hours IV or ORAL
[] famotidine (PEPCID) tablet	20 mg, oral, every 12 hours IV or ORAL
) pantoprazole (PROTONIX) IV or ORAL	"Or" Linked Panel
[] pantoprazole (PROTONIX) EC tablet	40 mg, oral, daily at 0600 Indication(s) for Proton Pump Inhibitor (PPI) Therapy:
 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:
() omeprazole (PriLOSEC) suspension	40 mg, Nasogastric, once, For 1 Doses Indication(s) for Proton Pump Inhibitor (PPI) Therapy:
Dexamethasone PO or IV (Single Response) -Dexamethasone should only be used in COVID- ventilator support. -Caution in using steroids early in COVID-19 dise	19 patients (a) requiring oxygen supplementation or (b) requiring pase (i.e. symptoms less than 7 days).
() devemethesens (DECADRON) tablet	6 mg arel deily For 10 Dagaa
 dexamethasone (DECADRON) tablet dexamethasone (DECADRON) IV 	6 mg, oral, daily, For 10 Doses 6 mg, intravenous, daily, For 10 Doses
() dexamethasone (DECADRON) IV	6 mg, oral, daily, For 10 Doses
	o mg, oral, dally, For To Doses
Antiemetics	
] ondansetron (ZOFRAN) IV	4 mg, intravenous, every 8 hours PRN, nausea, vomiting
] promethazine (PHENERGAN) IVPB or Oral or R	Rectal "Or" Linked Panel
[] promethazine (PHENERGAN) 25 mg in	12.5 mg, intravenous, for 30 Minutes, every 6 hours PRN, nausea,
sodium chloride 0.9 % 50 mL IVPB	vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required
[] promethazine (PHENERGAN) tablet	12.5 mg, oral, every 6 hours PRN, nausea, vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is able to tolera
	oral medication.
[] promethazine (PHENERGAN) suppository	12.5 mg, rectal, every 6 hours PRN, nausea, vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to
	tolerate oral medication.
Antiemetics	
] ondansetron (ZOFRAN) IV	4 mg, intravenous, every 8 hours PRN, nausea, vomiting
] promethazine (PHENERGAN) IV or Oral or Rect	
[] promethazine (PHENERGAN) 12.5 mg in	12.5 mg, intravenous, at 60 mL/hr, for 20 Minutes, every 6 hours PRN,
sodium chloride 0.9 % 0.9 % 20 mL for	nausea, vomiting
Alaris pump syringe option	Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to
	tolerate oral or rectal medication OR if a faster onset of action is require
[] promethazine (PHENERGAN) tablet	12.5 mg, oral, every 6 hours PRN, nausea, vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is able to tolera oral medication.
[] promethazine (PHENERGAN) suppository	12.5 mg, rectal, every 6 hours PRN, nausea, vomiting Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral medication.
Antiemetics	
] ondansetron (ZOFRAN) IV	4 mg, intravenous, every 8 hours PRN, nausea, vomiting
1 promethazine (PHENERGAN) IV or Oral or Rect	

[] promethazine (PHENERGAN) IV or Oral or Rectal

] promethazine (PHENERGAN) 12.5 mg IV	12.5 mg, intravenous, every 6 hours PRN, nausea, vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral or rectal medication OR if a faster onset of action is required.
[] promethazine (PHENERGAN) tablet	12.5 mg, oral, every 6 hours PRN, nausea, vomiting Give if ondansetron (ZOFRAN) is ineffective and patient is able to tolerate
[] promethazine (PHENERGAN) suppository	oral medication. 12.5 mg, rectal, every 6 hours PRN, nausea, vomiting
	Give if ondansetron (ZOFRAN) is ineffective and patient is UNable to tolerate oral medication.
Constipation (Single Response)	
() bisacodyl (DULCOLAX) EC tablet	10 mg, oral, daily PRN, constipation
() bisacodyl (DULCOLAX) suppository	10 mg, rectal, daily PRN, constipation
() lactulose solution	20 g, oral, every 8 hours PRN, constipation
() polyethylene glycol (GLYCOLAX) packet	17 g, oral, daily PRN, constipation
 () docusate (COLACE) 50 mg/5 mL liquid () docusate sodium (COLACE) capsule 	100 mg, Nasogastric, 2 times daily PRN, constipation 100 mg, oral, 2 times daily
Eye Care	
[] artificial tears ointment	Both Eyes, every 4 hours PRN, dry eyes Required for patients on paralytic agents or seventh cranial nerve palsy (Bell's Palsy)
[] hypromellose (NATURES TEARS) ophthalmic se	
Pain/Analgesia	
PRN Mild Pain (Pain Score 1-3) or Fever (Single	
Response)	
(adjust dose for renal/liver function and age)	
() acetaminophen (TYLENOL) tablet OR oral so	lution "Or" Linked Panel
Maximum of 3 grams of acetaminophen per d sources)	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from all
[] acetaminophen (TYLENOL) tablet	650 mg, oral, every 6 hours PRN, mild pain (score 1-3), fever, for fever GREATER than 102 F
	Maximum of 3 grams of acetaminophen per day from all sources.
[] acetaminophen (TYLENOL)suspension	(Cirrhosis patients maximum: 2 grams per day from all sources) 650 mg, oral, every 6 hours PRN, mild pain (score 1-3), fever, for fever
	GREATER than 102 F
	Maximum of 3 grams of acetaminophen per day from all sources.
	(Cirrhosis patients maximum: 2 grams per day from all sources). Use if patient cannot swallow tablet.
 PRN Oral Medications for Moderate Pain (Pain S 4-6): For Patients LESS than 65 years old (Single 	Score
Response) (adjust dose for renal/liver function and age)	
· · · · · · · · · · · · · · · · · · ·	
() acetaminophen-codeine (TYLENOL #3) table	
Maximum of 3 grams of acetaminophen per d sources)	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from all
[] acetaminophen-codeine (TYLENOL #3)	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6)
300-30 mg per tablet	Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if
[] acetaminophen-codeine 300 mg-30 mg /12.5 mL solution	 patient is able to tolerate oral medication. 12.5 mL, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if

	Maximum of 3 grams of acetaminophen per da	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from al
	sources)	
[]	HYDROcodone-acetaminophen (NORCO) 5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6)
[]	HYDROcodone-acetaminophen (HYCET) 2.5-108.3 mg/5 mL solution	10 mL, oral, every 6 hours PRN, moderate pain (score 4-6)
	HYDROcodone-acetaminophen 7.5/325 (NOR OR elixir	CO) tablet "Or" Linked Panel
	Maximum of 3 grams of acetaminophen per da sources)	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from all
[]	HYDROcodone-acetaminophen (NORCO) 7.5-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give i patient is able to tolerate oral medication.
[]	HYDROcodone-acetaminophen (HYCET) 7.5-325 mg/15 mL solution	15 mL, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient cannot swallow tablet.
	HYDROcodone-acetaminophen 10/325 (NOR) OR elixir	
	Maximum of 3 grams of acetaminophen per da sources)	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from al
[]	HYDROcodone-acetaminophen (NORCO 10-325) 10-325 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give patient is able to tolerate oral medication.
[]	HYDROcodone-acetaminophen (HYCET) 7.5-325 mg/15 mL solution	20 mL, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient can not swallow tablet.
()	traMADol (ULTRAM) tablet - For eGFR LESS than 30 mL/min, change frequency to every 12 hours)	50 mg, oral, every 6 hours PRN, moderate pain (score 4-6) (Max Daily dose not to exceed 200 mg/day). Give if patient is able to tolerate oral medication
4-6	RN Oral Medications for Moderate Pain (Pain S 6): For Patients GREATER than 65 years old (sponse)	core
	djust dose for renal/liver function and age)	
	acetaminophen-codeine (TYLENOL #3) tablet	
	Maximum of 3 grams of acetaminophen per da sources)	ay from all sources. (Cirrhosis patients maximum: 2 grams per day from all
[]	acetaminophen-codeine (TYLENOL #3) 300-30 mg per tablet	1 tablet, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources). Give if patient is able to tolerate oral medication.
[]	acetaminophen-codeine 300 mg-30 mg /12.5 mL solution	12.5 mL, oral, every 6 hours PRN, moderate pain (score 4-6) Maximum of 3 grams of acetaminophen per day from all sources. (Cirrhosis patients maximum: 2 grams per day from all sources) Use if patient cannot swallow tablet.
	HYDROcodone-acetaminophen 5/325 (NORC OR elixir	

() fentaNYL (SUB	LIMAZE) injection	50 mcg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
(adjust dose for re	nal/liver function and age)	
For Patients LESS	ns for Severe Pain (Pain Score 5 than 65 years old (Single Res	
release tablet	ROXICODONE) immediate	5 mg, oral, every 6 hours PRN, severe pain (score 7-10) Give if patient is able to tolerate oral medication.
() morphine (MSIF		15 mg, oral, every 6 hours PRN, severe pain (score 7-10) Give if patient is able to tolerate oral medication.
· · · ·	one (DILAUDID) tablet	2 mg, oral, every 6 hours PRN, severe pain (score 7-10) Give if patient is able to tolerate oral medication.
10-325) 10-325		1 tablet, oral, every 6 hours PRN, severe pain (score 7-10) Give if patient is able to tolerate oral medication.
7.5-325 mg per		1 tablet, oral, every 6 hours PRN, severe pain (score 7-10) Give if patient is able to tolerate oral medication.
Response) (adjust dose for re	nal/liver function and age)	
7-10): For Patients	s GREATER than 65 years old	
release tablet	ions for Severe Pain (Pain Sco	Give if patient is able to tolerate oral medication.
	ROXICODONE) immediate	Give if patient is able to tolerate oral medication. 10 mg, oral, every 6 hours PRN, severe pain (score 7-10)
() morphine (MSIF		Give if patient is able to tolerate oral medication. 15 mg, oral, every 6 hours PRN, severe pain (score 7-10)
	one (DILAUDID) tablet	2 mg, oral, every 6 hours PRN, severe pain (score 7-10)
Response)	s LESS than 65 years old (Sing nal/liver function and age)	
	ions for Severe Pain (Pain Sco	re
() HYDROmorpho	one (DILAUDID) injection	0.2 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
() morphine 2 mg/	/mL injection	1 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
() fentaNYL (SUB	LIMAZE) injection	12.5 mcg, intravenous, every 2 hour PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
	nal/liver function and age)	
	ns for Moderate Pain (Pain Sco ATER than 65 years old (Single	ore 4-6):
() HYDROmorpho	one (DILAUDID) injection	0.5 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
() morphine 2 mg/	/mL injection	2 mg, intravenous, every 3 hours PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
() fentaNYL (SUB	LIMAZE) injection	25 mcg, intravenous, every 2 hour PRN, moderate pain (score 4-6) Use if patient is unable to swallow or faster onset is needed
(adjust dose for re	nal/liver function and age)	
For Patients LESS	ns for Moderate Pain (Pain Sco 3 than 65 years old (Single Res	
every 12 hours)		Give if patient is able to tolerate oral medication
LESS than 30 n	RAM) tablet - For eGFR nL/min, change frequency to	25 mg, oral, every 6 hours PRN, moderate pain (score 4-6) (Max Daily dose not to exceed 200 mg/day).
2.5-108.3 mg/		10 mL, oral, every 6 hours PRN, moderate pain (score 4-6)
5-325 mg per t		

() morphine injection	4 mg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
() HYDROmorphone (DILAUDID) injection	0.8 mg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
] PRN IV Medications for Severe Pain (Pain Scor For Patients GREATER than 65 years old (Sing Response)	e 7-10):
(adjust dose for renal/liver function and age)	
() fentaNYL (SUBLIMAZE) injection	25 mcg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
() morphine injection	2 mg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
() HYDROmorphone (DILAUDID) injection	0.5 mg, intravenous, every 3 hours PRN, severe pain (score 7-10) Use if patient is unable to swallow or faster onset is needed
nsomnia	
] ramelteon (ROZEREM) tablet	8 mg, oral, nightly PRN, sleep
Respiratory Inhalers	
] albuterol (PROAIR HFA) inhaler	2 puff, inhalation, every 4 hours PRN, wheezing MDI with spacer only
] ipratropium (ATROVENT HFA) inhaler	2 puff, inhalation, every 4 hours PRN, wheezing, shortness of breath MDI with spacer only
sodium chloride 0.9% bag for line care	
[X] sodium chloride 0.9% bag for line care	250 mL, intravenous, PRN, line care For flushing of extension tubing sets after administration of intermittent infusions. Program sodium chloride bag to run at the same infusion rate as medication given for a total volume equal to contents of tubing sets used. Change bag every 24 hours.
VTE	
VIE DVT Risk and Prophylaxis Tool (Single Respons	(Selection Required)
	URL: "\appt1.pdf"
() Patient currently has an active order for therape anticoagulant or VTE prophylaxis	eutic Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
() LOW Risk of DVT (Selection Required)	
Low Risk Definition Age less than 60 years and NO other VTE risk f	actors
[] Low Risk (Single Response) (Selection Requ	ired)
() Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourga early ambulation
() MODERATE Risk of DVT - Surgical (Selection F	

contraindicated. One or more of the following medical conditions:	lechanical prophylaxis is optional unless pharmacologic is
	nation, dehydration, varicose veins, cancer, sepsis, obesity, previous leg swelling, ulcers, venous stasis and nephrotic syndrome
History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hour Less than fully and independently ambulatory	rs
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 - S Patient (Single Response) (Selection Required) 	Surgical
() Contraindications exist for pharmacologic prop BUT order Sequential compression device	
 [] 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 prop AND mechanical prophylaxis	
 [] 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 Resp (Selection Required) 	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() 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
() 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
() patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30 mL/min
() fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1 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.
() warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:

()	Pharmacy consult to manage warfarin
	(COUMADIN)

() MODERATE Risk of DVT - Non-Surgical (Selectio Required)	n	
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, previ 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 hour Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission	S	
Major surgery within 5 months of admission		
 [] Moderate Risk (Selection Required) [] Moderate risk of VTE [] Moderate Risk Pharmacological Prophylaxis - 	Routine, Once	
Non-Surgical Patient (Single Response) (Select Required)	tion	
 () Contraindications exist for pharmacologic prop Order Sequential compression device 	ohylaxis - "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 prop AND mechanical prophylaxis	ohylaxis "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 Resp (Selection Required)		
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S	
() 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	
() 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	
() patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min	
() fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily 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	
 () 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.	
() warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:	

() Pharmacy consult to manage warfarin	STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
() HIGH Risk of DVT - Surgical (Selection Required)	
High Risk Definition	and the end decreased
Both pharmacologic AND mechanical prophylaxis	must be addressed.
One or more of the following medical conditions:	ant mutations, anticardiolipin antibody syndrome; antithrombin, protein C
or protein S deficiency; hyperhomocysteinemia; m	
Severe fracture of hip, pelvis or leg	ryeloprometative disorders/
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)	Douting Ones
[] High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Surgi (Single Response) (Selection Required)	
() Contraindications exist for pharmacologic	Routine, Once
prophylaxis	No pharmacologic VTE prophylaxis due to the following
	contraindication(s):
() enoxaparin (LOVENOX) injection (Single Res (Selection Required)	ponse)
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() 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
() patients weight between 100-139 kg AND	30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
CrCl GREATER than 30 mL/min	For Patients weight between 100-139 kg and CrCl GREATER than 30
	mL/min
() patients weight 140 kg or GREATER AND	40 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
CrCl GREATER than 30 mL/min	For Patients weight 140 kg or GREATER and CrCl GREATER than 30
	mL/min
() fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1
	If the patient does not have a history or suspected case of
	Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication.
	Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCl LESS than 30 mL/min.
	This patient has a history of or suspected case of Heparin-Induced
	Thrombocytopenia (HIT):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM
for patients with high risk of bleeding, e.g.	Recommended for patients with high risk of bleeding, e.g. weight LESS
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() warf arin (COUMADIN) tablet	oral, daily at 1700, Starting S+1
	Indication:
() Pharmacy consult to manage warfarin	STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
[] Mechanical Prophylaxis (Single Response) (Se	lection
Required)	
() Contraindications exist for mechanical	Routine, Once
prophylaxis	No mechanical VTE prophylaxis due to the following contraindication(s):
	No mechanical VTE prophylaxis due to the following contraindication(s): Routine, Continuous

High Risk Definition Both pharmacologic AND mechanical prophylax One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin va or protein S deficiency; hyperhomocysteinemia; 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	: riant mutations, anticardiolipin antibody syndrome; antithrombin, protein C
[] High Risk (Selection Required)	
[] High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Nor Patient (Single Response) (Selection Require	
() Contraindications exist for pharmacologic	Routine, Once
prophylaxis	No pharmacologic VTE prophylaxis due to the following
	contraindication(s):
 enoxaparin (LOVENOX) injection (Single Re (Selection Required) 	sponse)
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S
() 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
() 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
() patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min
() fondaparinux (ARIXTRA) injection	 2.5 mg, subcutaneous, daily 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
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours
for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs)	Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
() warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
[] Mechanical Prophylaxis (Single Response) (S Required)	Selection
() 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 of DVT - Surgical (Hip/Knee) (Select	ion

() HIGH Risk Required)

High Risk Definition Both pharmacologic AND mechanical prophylaxis One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin vari	s must be addressed. ant mutations, anticardiolipin antibody syndrome; antithrombin, protein C
or protein S deficiency; hyperhomocysteinemia; n	
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)	
[] High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Hip of	
(Arthroplasty) Surgical Patient (Single Respon	
(Selection Required)	
() Contraindications exist for pharmacologic	Routine, Once
prophylaxis	No pharmacologic VTE prophylaxis due to the following
propriylaxio	contraindication(s):
() apixaban (ELIQUIS) tablet	2.5 mg, oral, every 12 hours, Starting S+1
	Indications:
() aspirin chewable tablet	162 mg, oral, daily, Starting S+1
() aspirin (ECOTRIN) enteric coated tablet	162 mg, oral, daily, Starting S+1
() enoxaparin (LOVENOX) injection (Single Res	
() enotapain (LOVENOX) injection (Single Res (Selection Required)	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() enoxaparin (LOVENOX) syringe	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL),
() chokapalin (Eovervox) synnge	Starting S+1
() enoxaparin (LOVENOX) syringe - For	30 mg, subcutaneous, daily at 0600, Starting S+1
Patients with CrCL LESS than 30 mL/min	For Patients with CrCL LESS than 30 mL/min.
() enoxaparin (LOVENOX) syringe - For	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL),
Patients weight between 100-139 kg and	Starting S+1
CrCl GREATER than 30 mL/min	For Patients weight between 100-139 kg and CrCl GREATER than 30
	mL/min.
() enoxaparin (LOVENOX) syringe - For	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL),
Patients weight between 140 kg or	Starting S+1
GREATER and CrCI GREATER than 30	For Patients weight 140 kg or GREATER and CrCl GREATER than 30
mL/min	mL/min
() 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):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM
for patients with high risk of bleeding, e.g.	Recommended for patients with high risk of bleeding, e.g. weight LESS
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() rivaroxaban (XARELTO) tablet for hip or	10 mg, oral, daily at 0600 (TIME CRITICAL), Starting S+1
knee arthroplasty planned during this	To be Given on Post Op Day 1.
admission	Indications:
() warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1
· · ·	Indication:
() Pharmacy consult to manage warfarin	STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
[] Mechanical Prophylaxis (Single Response) (Se	election
Required)	
() Contraindications exist for mechanical	Routine, Once
prophylaxis	No mechanical VTE prophylaxis due to the following contraindication(s):

() Place/Maintain sequential compression device continuous	Routine, Continuous
DVT Risk and Prophylaxis Tool (Single Response)	(Selection Required) URL: "\appt1.pdf"
() Patient currently has an active order for therapeut anticoagulant or VTE prophylaxis	ic Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
() LOW Risk of DVT (Selection Required)	
Low Risk Definition Age less than 60 years and NO other VTE risk fac	tors
[] Low Risk (Single Response) (Selection Require	ed)
() Low risk of VTE	Routine, Once Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourgae early ambulation
() MODERATE Risk of DVT - Surgical (Selection Re	
Moderate Risk Definition	····/
Pharmacologic prophylaxis must be addressed. M	lechanical prophylaxis is optional unless pharmacologic is
contraindicated.	
stroke, rheumatologic disease, sickle cell disease, Age 60 and above	nation, dehydration, varicose veins, cancer, sepsis, obesity, previous leg swelling, ulcers, venous stasis and nephrotic syndrome
Central line	
History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hour	29
Less than fully and independently ambulatory	3
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 - S Patient (Single Response) (Selection Required)	Surgical
() Contraindications exist for pharmacologic prop BUT order Sequential compression device	-
[] 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 prop AND mechanical prophylaxis 	-
[] 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 Resp (Selection Required)	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() 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
() 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
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() 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
() 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.
() warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
[] Mechanical Prophylaxis (Single Response) (Se Required)	lection
 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 of DVT - Non-Surgical (Selection Required)	n
Age 60 and above Central line History of DVT or family history of VTE Anticipated length of stay GREATER than 48 hour Less than fully and independently ambulatory Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission	rs
[] Moderate Risk (Selection Required)	
 [] Moderate risk of VTE [] Moderate Risk Pharmacological Prophylaxis - Non-Surgical Patient (Single Response) (Selec Required) () Contraindications exist for pharmacologic prop 	
Order Sequential compression device	ohvlaxis - "And" Linked Panel
[] Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
[] Contraindications exist for pharmacologic	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): Routine, Continuous
 [] Contraindications exist for pharmacologic prophylaxis [] Place/Maintain sequential compression device continuous () Contraindications exist for pharmacologic properties of the second s	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): Routine, Continuous
 [] Contraindications exist for pharmacologic prophylaxis [] Place/Maintain sequential compression device continuous 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): Routine, Continuous

() enoxaparin (LOVENOX) injection (Single Re (Selection Required)	· · ·
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S+1
() 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
() 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 3 mL/min
() 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 3 mL/min
() 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):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours
 () heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs) () warfarin (COUMADIN) tablet 	5,000 Units, subcutaneous, every 12 hours Recommended for patients with high risk of bleeding, e.g. weight LES than 50kg and age GREATER than 75yrs. oral, daily at 1700
() Pharmacy consult to manage warfarin	Indication: STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
HIGH Risk of DVT - Surgical (Selection Require	d)
Address both pharmacologic and mechanical pro	ophylaxis by ordering from Pharmacological and Mechanical Prophylaxis.
] High Risk (Selection Required)	
[] High risk of VTE	Routine, Once
] High Risk Pharmacological Prophylaxis - Surg	gical Patient
(Single Response) (Selection Required)	
	Routine, Once No pharmacologic VTE prophylaxis due to the following
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Response) 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Reguired) 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse)
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Response) 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Re (Selection Required) () enoxaparin (LOVENOX) syringe 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min 30 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Re(Selection Required) () enoxaparin (LOVENOX) syringe () patients with CrCL LESS than 30 mL/min () patients weight between 100-139 kg AND 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS 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 3 mL/min 40 mg, subcutaneous, 2 times daily at 0600, 1800, Starting S+1
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Re(Selection Required) () enoxaparin (LOVENOX) syringe () patients with CrCL LESS than 30 mL/min () patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min () patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min () fondaparinux (ARIXTRA) injection 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS 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 3 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 3 mL/min 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 medicati 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):
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Reservent) () enoxaparin (LOVENOX) syringe () enoxaparin (LOVENOX) syringe () patients with CrCL LESS than 30 mL/min () patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min () patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min () fondaparinux (ARIXTRA) injection 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS 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 3 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 3 mL/min 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 medicati 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): 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Reserver) () enoxaparin (LOVENOX) syringe () patients with CrCL LESS than 30 mL/min () patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min () patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min () fondaparinux (ARIXTRA) injection () heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS 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 3 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 3 mL/min 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 medicati 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): 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM 5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM
 (Single Response) (Selection Required) () Contraindications exist for pharmacologic prophylaxis () enoxaparin (LOVENOX) injection (Single Re(Selection Required) () enoxaparin (LOVENOX) syringe () patients with CrCL LESS than 30 mL/min () patients weight between 100-139 kg AND CrCl GREATER than 30 mL/min () patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min () fondaparinux (ARIXTRA) injection () heparin (porcine) injection (Recommended 	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): esponse) 40 mg, subcutaneous, daily at 0600, Starting S+1 30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS 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 3 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 3 mL/min 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 medicati 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): 5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM

) HIGH Risk of DVT - Non-Surgical (Selection Requ	uired)
	phylaxis by ordering from Pharmacological and Mechanical Prophylaxis.
[] High Risk (Selection Required)	
[] High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Non- Patient (Single Response) (Selection Required	
() Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
 enoxaparin (LOVENOX) injection (Single Res (Selection Required) 	ponse)
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily, Starting S+1
() patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily, Starting S+1 For Patients with CrCL LESS than 30 mL/min
() 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
() 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
() 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):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours
for patients with high risk of bleeding, e.g.	Recommended for patients with high risk of bleeding, e.g. weight LESS
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
 Pharmacy consult to manage warfarin (COUMADIN) 	STAT, Until discontinued, Starting S Indication:
) HIGH Risk of DVT - Surgical (Hip/Knee) (Selectio	n
Required)	phylaxis by ordering from Pharmacological and Mechanical Prophylaxis.
Address both pharmacologic and mechanical pro	
[] High Risk (Selection Required)	
[] High risk of VTE	Routine, Once
[] High Risk Pharmacological Prophylaxis - Hip o	rKnee
(Arthroplasty) Surgical Patient (Single Respons (Selection Required)	se)
() Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
() apixaban (ELIQUIS) tablet	2.5 mg, oral, every 12 hours, Starting S+1 Indications:
() aspirin chewable tablet	162 mg, oral, daily, Starting S+1
() aspirin (ECOTRIN) enteric coated tablet	162 mg, oral, daily, Starting S+1
 enoxaparin (LOVENOX) injection (Single Res (Selection Required) 	ponse)
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() enoxaparin (LOVENOX) syringe	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1
() enoxaparin (LOVENOX) syringe - For Patients with CrCL LESS than 30 mL/min	30 mg, subcutaneous, daily at 0600, Starting S+1 For Patients with CrCL LESS than 30 mL/min.

() enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1
CrCl GREATER than 30 mL/min	For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min.
() enoxaparin (LOVENOX) syringe - For	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL),
Patients weight between 140 kg or GREATER and CrCl GREATER than 30	Starting S+1 For Patients weight 140 kg or GREATER and CrCI GREATER than 30
mL/min	mL/min
() 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 medicatior Contraindicated in patients LESS than 50kg, prior to surgery/invasive procedure, or CrCI LESS than 30 mL/min
	This patient has a history of or suspected case of Heparin-Induced
	Thrombocytopenia (HIT):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
 heparin (porcine) injection (Recommended for patients with high risk of bleeding, e.g. 	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
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() rivaroxaban (XARELTO) tablet for hip or	10 mg, oral, daily at 0600 (TIME CRITICAL), Starting S+1
knee arthroplasty planned during this	To be Given on Post Op Day 1.
admission	Indications:
() warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1
() Pharmacy consult to manage warfarin	Indication: STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
VT Risk and Prophylaxis Tool (Single Response)) URL: "\appt1.pdf"
) Patient currently has an active order for therapeut anticoagulant or VTE prophylaxis	ic Routine, Once No pharmacologic VTE prophylaxis because: patient is already on therapeutic anticoagulation for other indication. Therapy for the following:
) LOW Risk of DVT (Selection Required)	
Low Risk Definition	
Age less than 60 years and NO other VTE risk fac	ctors
[] Low Risk (Single Response) (Selection Require	2d)
() Low risk of VTE	Routine, Once
	Low risk: Due to low risk, no VTE prophylaxis is needed. Will encourgae
	early ambulation
MODERATE Risk of DVT - Surgical (Selection Re	quired)
Moderate Risk Definition Pharmacologic prophylaxis must be addressed. M contraindicated.	lechanical prophylaxis is optional unless pharmacologic is
One or more of the following medical conditions:	
	nation, dehydration, varicose veins, cancer, sepsis, obesity, previous , leg swelling, ulcers, venous stasis and nephrotic syndrome
Age 60 and above	, ובט איפווווש, טונפוס, יפווטטס סנמסוס מווט וופטווטנוג סטוטוטוופ
Central line	
History of DVT or family history of VTE	
Anticipated length of stay GREATER than 48 hou	rs
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 - Patient (Single Response) (Selection Required	
 Contraindications exist for pharmacologic pro BUT order Sequential compression device 	
[] 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 pro AND mechanical prophylaxis	phylaxis "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 Res (Selection Required)	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() 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
() 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
() patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1 For Patient weight of 140 kg or GREATER and CrCl GREATER than 30
	mL/min
() fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1
	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.	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
weight < 50kg and age > 75yrs) () warf arin (COUMADIN) tablet	than 50kg and age GREATER than 75yrs. oral, daily at 1700, Starting S+1
() warfarin (COUMADIN) tablet	Indication:
() Pharmacy consult to manage warfarin	STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
() MODERATE Risk of DVT - Non-Surgical (Selection	on
Required) Moderate Risk Definition	
	Mechanical prophylaxis is optional unless pharmacologic is
contraindicated.	viechanical propriyaxis is optional unless pharmacologic is
One or more of the following medical conditions:	
	mation, dehydration, varicose veins, cancer, sepsis, obesity, previous e, leg swelling, ulcers, venous stasis and nephrotic syndrome
Central line History of DVT or family history of VTE	
Anticipated length of stay GREATER than 48 hou Less than fully and independently ambulatory	Irs
Estrogen therapy Moderate or major surgery (not for cancer) Major surgery within 3 months of admission	
1 Moderate Disk (Selection Required)	

[] Moderate risk of VTE	Routine, Once
[] Moderate Risk Pharmacological Prophylaxis -	
Non-Surgical Patient (Single Response) (Selec	tion
Required)	shulovia "And" Linked Denel
() Contraindications exist for pharmacologic prop Order Sequential compression device	-
 [] 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 prop AND mechanical prophylaxis	ohylaxis "And" Linked Panel
[] Contraindications exist for pharmacologic prophylaxis	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
[] Contraindications exist for mechanical	Routine, Once
prophylaxis	No mechanical VTE prophylaxis due to the following contraindication(s):
() enoxaparin (LOVENOX) injection (Single Resp (Selection Required)	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 1700, Starting S
() 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
() patients weight between 100-139 kg AND	30 mg, subcutaneous, 2 times daily, Starting S
CrCl GREATER than 30 mL/min	For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min
() patients weight 140 kg or GREATER AND	40 mg, subcutaneous, 2 times daily, Starting S
CrCl GREATER than 30 mL/min	For Patients weight 140 kg or GREATER and CrCl GREATER than 3 mL/min
() 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):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours
for patients with high risk of bleeding, e.g. weight < 50kg and age > 75yrs)	Recommended for patients with high risk of bleeding, e.g. weight LESS than 50kg and age GREATER than 75yrs.
() warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
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:	
	ant mutations, anticardiolipin antibody syndrome; antithrombin, protein C
or protein S deficiency; hyperhomocysteinemia; m	yeloproliferative 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)	
[] High risk of VTE	Routine, Once

	Routine, Once
() Contraindications exist for pharmacologic prophylaxis	No pharmacologic VTE prophylaxis due to the following contraindication(s):
 enoxaparin (LOVENOX) injection (Single Resp (Selection Required) 	
() enoxaparin (LOVENOX) syringe	40 mg, subcutaneous, daily at 0600, Starting S+1
() 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
() 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
() 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
() 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):
() 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.	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
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() warfarin (COUMADIN) tablet	oral, daily at 1700, Starting S+1 Indication:
() Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
[] Mechanical Prophylaxis (Single Response) (Sel Required)	
 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 of DVT - Non-Surgical (Selection Requ	irea)
High Risk Definition Both pharmacologic AND mechanical prophylaxis One or more of the following medical conditions: Thrombophilia (Factor V Leiden, prothrombin varia or protein S deficiency; hyperhomocysteinemia; m	nt mutations, anticardiolipin antibody syndrome; antithrombin, protein C
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)	Dauting Ones
[] High risk of VTE [] High Risk Pharmacological Prophylaxis - Non-S	Routine, Once Surgical
Patient (Single Response) (Selection Required)	Poutino Onco
() Contraindications exist for pharmacologic	Routine, Once No pharmacologic VTE prophylaxis due to the following
prophylaxis	contraindication(s):

()		
	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
()	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
()	patients weight 140 kg or GREATER AND CrCl GREATER than 30 mL/min	40 mg, subcutaneous, 2 times daily, Starting S For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min
()	fondaparinux (ARIXTRA) injection	 2.5 mg, subcutaneous, daily 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
()	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.
()	warfarin (COUMADIN) tablet	oral, daily at 1700 Indication:
()	Pharmacy consult to manage warfarin (COUMADIN)	STAT, Until discontinued, Starting S Indication:
	Mechanical Prophylaxis (Single Response) (Se Required)	
()	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
HIG	GH Risk of DVT - Surgical (Hip/Knee) (Selection	1
	quired)	
One Thre or p Sev	protein S deficiency; hyperhomocysteinemia; m vere fracture of hip, pelvis or leg	ant mutations, anticardiolipin antibody syndrome; antithrombin, protein C
Mu Abo Acu	cute spinal cord injury with paresis Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke tory of PE	
Mul Abo Acu His	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required)	
Mul Abo Acu His	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke tory of PE High Risk (Selection Required) High risk of VTE	Routine, Once
Mul Abo Acu His [] H [] H	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required)	rKnee
Mul Abo His [] H [] H [] H ((Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke tory of PE High Risk (Selection Required) High risk of VTE High Risk Pharmacological Prophylaxis - Hip or (Arthroplasty) Surgical Patient (Single Respons	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s):
Mul Abo His [] H [] [] (()	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required) High risk of VTE High Risk Pharmacological Prophylaxis - Hip or (Arthroplasty) Surgical Patient (Single Respons (Selection Required) Contraindications exist for pharmacologic prophylaxis apixaban (ELIQUIS) tablet	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): 2.5 mg, oral, every 12 hours, Starting S+1 Indications:
Mul Abo His [] H [] H (()	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required) High Risk of VTE High Risk Pharmacological Prophylaxis - Hip or (Arthroplasty) Surgical Patient (Single Respons (Selection Required) Contraindications exist for pharmacologic prophylaxis apixaban (ELIQUIS) tablet aspirin chewable tablet	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): 2.5 mg, oral, every 12 hours, Starting S+1 Indications: 162 mg, oral, daily, Starting S+1
Mul Abo Acu His [] F [] F ((() () ()	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required) High risk of VTE High Risk Pharmacological Prophylaxis - Hip or (Arthroplasty) Surgical Patient (Single Respons (Selection Required) Contraindications exist for pharmacologic prophylaxis apixaban (ELIQUIS) tablet aspirin chewable tablet aspirin (ECOTRIN) enteric coated tablet enoxaparin (LOVENOX) injection (Single Resp	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): 2.5 mg, oral, every 12 hours, Starting S+1 Indications: 162 mg, oral, daily, Starting S+1 162 mg, oral, daily, Starting S+1
Mul Abo Acu His [] F [] F ((() () ()	Itiple major traumas dominal or pelvic surgery for CANCER ute ischemic stroke story of PE High Risk (Selection Required) High risk of VTE High Risk Pharmacological Prophylaxis - Hip or (Arthroplasty) Surgical Patient (Single Respons (Selection Required) Contraindications exist for pharmacologic prophylaxis apixaban (ELIQUIS) tablet aspirin chewable tablet aspirin (ECOTRIN) enteric coated tablet	Routine, Once No pharmacologic VTE prophylaxis due to the following contraindication(s): 2.5 mg, oral, every 12 hours, Starting S+1 Indications: 162 mg, oral, daily, Starting S+1 162 mg, oral, daily, Starting S+1

(

() 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.
() enoxaparin (LOVENOX) syringe - For Patients weight between 100-139 kg and	30 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1
CrCl GREATER than 30 mL/min	For Patients weight between 100-139 kg and CrCl GREATER than 30 mL/min.
() enoxaparin (LOVENOX) syringe - For Patients weight between 140 kg or	40 mg, subcutaneous, 2 times daily at 0600, 1800 (TIME CRITICAL), Starting S+1
GREATER and CrCl GREATER than 30 mL/min	For Patients weight 140 kg or GREATER and CrCl GREATER than 30 mL/min
() fondaparinux (ARIXTRA) injection	2.5 mg, subcutaneous, daily, Starting S+1
	If the patient does not have a history or suspected case of
	Heparin-Induced Thrombocytopenia (HIT) do NOT order this medication
	Contraindicated in patients LESS than 50kg, prior to surgery/invasive
	procedure, or CrCl LESS than 30 mL/min This patient has a history of or suspected case of Heparin-Induced
	Thrombocytopenia (HIT):
() heparin (porcine) injection	5,000 Units, subcutaneous, every 8 hours, S+1 at 6:00 AM
() heparin (porcine) injection (Recommended	5,000 Units, subcutaneous, every 12 hours, S+1 at 6:00 AM
for patients with high risk of bleeding, e.g.	Recommended for patients with high risk of bleeding, e.g. weight LESS
weight < 50kg and age > 75yrs)	than 50kg and age GREATER than 75yrs.
() rivaroxaban (XARELTO) tablet for hip or knee arthroplasty planned during this	10 mg, oral, daily at 0600 (TIME CRITICAL), Starting S+1 To be Given on Post Op Day 1.
admission	Indications:
() warf arin (COUMADIN) tablet	oral, daily at 1700, Starting S+1
	Indication:
() Pharmacy consult to manage warfarin	STAT, Until discontinued, Starting S
(COUMADIN)	Indication:
] Mechanical Prophylaxis (Single Response) (Se	election
Required)	
() Contraindications exist for mechanical	Routine, Once
prophylaxis () Place/Maintain sequential compression	No mechanical VTE prophylaxis due to the following contraindication(s):
() Place/Maintain sequential compression device continuous	Routine, Continuous

Labs

Laboratory-Admission

[X] CBC with platelet and differential	STAT For 1 Occurrences
[X] Comprehensive metabolic panel	STAT For 1 Occurrences
[X] Prothrombin time with INR	STAT For 1 Occurrences
[X] Partial thromboplastin time, activated (PTT)	STAT For 1 Occurrences
[X] Troponin	STAT For 1 Occurrences
[X] BNP	STAT For 1 Occurrences
[X] Myoglobin	STAT For 1 Occurrences
[X] Procalcitonin	STAT For 1 Occurrences
[X] IgG subclasses	STAT For 1 Occurrences
[X] Creatine kinase, total (CPK)	STAT For 1 Occurrences
[] hCG qualitative, urine screen	STAT For 1 Occurrences

Laboratory-Inflammatory Bundle

[X] C-reactive protein	Once
[X] Interleukin 6	Once
[X] Ferritin level	Once
[X] D-dimer	Once
[X] LDH	Once
[X] Triglycerides	Once
[X] Fibrinogen	Once
[] Prothrombin time with INR	Once
[] Partial thromboplastin time, activated	Once

[X] CBC with platelet and differential	AM draw repeats For 3 Occurrences
[X] Comprehensive metabolic panel	AM draw repeats For 3 Occurrences
[] Additional Daily labs-Critical Illness/Clinical De	
Consider these daily repeat labs with Moderat	te/Severe IIIness in COVID-19 positive patients.
[] Troponin	AM draw repeats, Starting S+1 For 3 Occurrences
[] D-dimer	AM draw repeats, Starting S+1 For 3 Occurrences
[] C-reactive protein	AM draw repeats, Starting S+1 For 3 Occurrences
[] LDH	AM draw repeats, Starting S+1 For 3 Occurrences
[] Ferritin level	AM draw repeats, Starting S+1 For 3 Occurrences
Laboratory-Type and Screen	
[X] Type and screen	STAT For 1 Occurrences
Cardiology	
Cardiology	
[X] ECG 12 lead upon admission	Routine, STAT For 1 Occurrences
	Clinical Indications: Rate/Rhythm
	Interpreting Physician:
[] ECG 12 lead daily	Routine, Daily For 3 Occurrences
	Clinical Indications: Interpreting Physician:
[] Transthoracic Echocardiogram Complete, (w Strain and 3D if needed)	
Imaging	
CXR	
[X] XR Chest 1 Vw Portable	
	STAT, 1 time imaging For 1 Occurrences
[] Daily XR Chest 1 Vw Portable	Routine, Daily imaging, Starting S+1 For Until specified
	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI >
Daily XR Chest 1 Vw Portable	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor.
Daily XR Chest 1 Vw Portable	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI >
Daily XR Chest 1 Vw Portable	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine
Daily XR Chest 1 Vw Portable Respiratory Respiratory	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI = 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation:
Daily XR Chest 1 Vw Portable Respiratory Respiratory	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI = 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies:
Daily XR Chest 1 Vw Portable Respiratory Respiratory	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies:
Daily XR Chest 1 Vw Portable Respiratory Respiratory	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies:
Daily XR Chest 1 Vw Portable Respiratory Respiratory [] Mechanical ventilation	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies:
Daily XR Chest 1 Vw Portable Respiratory Respiratory [] Mechanical ventilation	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI : 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies:
Daily XR Chest 1 Vw Portable Respiratory Respiratory [] Mechanical ventilation	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI : 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Routine, Continuous Device: Titrate to keep O2 Sat Above:
 Daily XR Chest 1 Vw Portable Respiratory Respiratory Mechanical ventilation 	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Routine, Continuous Device: Titrate to keep O2 Sat Above: Indications for O2 therapy:
 Daily XR Chest 1 Vw Portable Respiratory Respiratory Mechanical ventilation 	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Routine, Continuous Device: Titrate to keep O2 Sat Above: Indications for O2 therapy: Device 2:
 Daily XR Chest 1 Vw Portable Respiratory Respiratory Mechanical ventilation 	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI : 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Routine, Continuous Device: Titrate to keep O2 Sat Above: Indications for O2 therapy: Device 2: Device 3:
[] Daily XR Chest 1 Vw Portable Respiratory [] Mechanical ventilation [] Oxygen therapy-	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Device: Titrate to keep O2 Sat Above: Indications for O2 therapy: Device 3: Indications for O2 therapy:
Daily XR Chest 1 Vw Portable Respiratory Respiratory [] Mechanical ventilation	Routine, Daily imaging, Starting S+1 For Until specified Consider daily CXR for the following patients: Age > 70, BMI > 40, or Increasing O2 requirements on the floor. URL: "\appt1Hypoxemia Algorithm.pdf" Routine Mechanical Ventilation: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Vent Management Strategies: Routine, Continuous Device: Titrate to keep O2 Sat Above: Indications for O2 therapy: Device 2: Device 3:

Physician Consults Consider using these consults to assist with management of the COVID-19 positive patient.

[]	Consult Infectious Diseases for moderate to severe COVID-19 patient	Reason for Consult? Management of COVID-19 positive patient
		Patient/clinical information communicated?
[]	Consult Hematology and Oncology for suspected	Reason for Consult? Management of COVID-19 positive
	Cytokine Storm	patient with suspected Cytokine Storm
		Patient/clinical information communicated?
[]	Consult Pulmonary/Crit Care for respiratory insufficiency	Reason for Consult? Management of COVID-19 positive
	patient with respiratory insufficiency	
		Patient/clinical information communicated?
[]	Consult Nephrology/Hyperten	Reason for Consult?
		Patient/Clinical information communicated?
		Patient/clinical information communicated?
Ar	ncillary Consults	
	cillary Consults	
An	cillary Consults	
		Priority: Same Day
An	cillary Consults	
An	cillary Consults	Priority: Same Day Reason for Consult? Assistance with clarification of goals of
An	cillary Consults	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care
An	cillary Consults	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care Order?
A n	cillary Consults	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care Order? Name of referring provider:
An	cillary Consults Consult to Palliative Care Service	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care Order? Name of referring provider: Enter call back number:
An	cillary Consults Consult to Palliative Care Service	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care Order? Name of referring provider: Enter call back number: Reason For Consult?
A n	cillary Consults Consult to Palliative Care Service Consult to Nutrition Services	Priority: Same Day Reason for Consult? Assistance with clarification of goals of care Order? Name of referring provider: Enter call back number: Reason For Consult? Purpose/Topic: