





THE UNIVERSITY  
OF KANSAS HOSPITAL  
**KUMED**

3901 Rainbow Boulevard  
Kansas City, Kansas 66160

**PHYSICIAN'S ORDER FORM**

PATIENT LABEL

Room

DATE & TIME	#	ORDERS ADULT SEVERE SEPSIS/ SEPTIC SHOCK ORDER SET
	10.	<p>OPTIMIZATION OF MAP (mean arterial pressure)</p> <p>a. Check or calculate MAP (MAP = [2 X DBP] + SBP/ 3)</p> <p>b. If MAP is less than 65 mmHg, give vasopressor of choice to maintain a MAP <math>\geq</math> 65 mmHg.  <input type="checkbox"/> Norepinephrine _____ mcg/min (range 2-30 mcg/min)  <input type="checkbox"/> Vasopressin _____ units/min (range 0.01-0.06 units/min)  <input type="checkbox"/> Epinephrine _____ mcg/min (range 2-10 mcg/min)</p> <p>c. When goal MAP of <math>\geq</math> 65 mmHg is achieved, continue to OPTIMIZATION OF ScvO2.</p>
	11.	<p>OPTIMIZATION OF ScvO2 (Mixed central venous O2 sat)</p> <p>a. Check ScvO2.</p> <p>b. If ScvO2 is less than 70% and Hbg is less than 10 grams, transfuse _____ units packed red blood cells until Hbg is <math>\geq</math> 10 grams and recheck ScvO2.</p> <p>c. Dobutamine 2.5mcg/kg/min infuse, if ScvO2 is less than 70% and Hbg is <math>\geq</math> 10 grams, increase every 30 minutes until ScvO2 is at least 70%. Maximum dose of dobutamine not to exceed 20 mcg/kg/min. (Usual dose 2.5-10 mcg/kg/min)</p> <p>d. If ScvO2 is <math>\geq</math> 70%, early goal directed therapy is complete.</p> <p>*** REASSESS EACH STEP EVERY 30 MINUTES TO MAINTAIN OPTIMIZATION GOALS.***</p>
	12.	Ventilation – follow “Acute Lung Injury Protocol” if mechanically ventilated.
	13.	<p>Steroids: (select all that all applicable)</p> <p><input type="checkbox"/> Hydrocortisone IV 100mg every 8 hours once cortisol level is drawn. <b>(Cortisol level must be ordered)</b>            (If cortisol level is greater than 25mcg/dL, please write a separate order on the physician order sheet to discontinue IV hydrocortisone <b>and make a note of the cortisol level after you order.</b>)</p> <p><input type="checkbox"/> Continue hydrocortisone for 7 days at the above dose OR</p> <p><input type="checkbox"/> Discontinue hydrocortisone when the patient no longer has a vasopressor requirement.</p> <p><input type="checkbox"/> Fludrocortisone 0.05mg daily PO/NG along with IV hydrocortisone.</p> <p><b>Reason not given:</b></p> <p><input type="checkbox"/> Patient not on vasopressors</p> <p><input type="checkbox"/> MAP, CVP, Scvo2 (Svo2) goals met</p> <p><input type="checkbox"/> Cortisol level within normal limits</p> <p><input type="checkbox"/> Other (please list reason):</p>
	14.	Once Early Goal Directed Therapy goals are achieved, recheck lactate every 4 hours until normal. ( $\leq$ 2 mmol/L).
	15.	Use the “Critical Care Intensive IV Insulin Infusion Protocol” order set for tight glucose control.
	16.	Initiate ICU based admission order sets for DVT and Stress Ulcer Prophylaxis.
	17.	<p>Notify physician if:</p> <p>a. CVP &lt; 8 or &gt; 15 mmHg</p> <p>b. ScvO2 &lt; 70%</p> <p>c. Hemoglobin &lt; 10g/dL</p> <p>d. If lactate is &gt; 2 mmol/L</p> <p>e. If oxygen saturation &lt; 88% or inspiratory plateau &gt; 30 cm H<sub>2</sub>O (on mechanical ventilation)</p>

Physician Signature: \_\_\_\_\_ Pager: \_\_\_\_\_

**ADULT SEVERE SEPSIS/ SEPTIC SHOCK ORDER SET**

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Room

DATE & TIME	#	ORDERS
		<b>ACTIVATED PROTEIN C / DROTRECOGIN ALFA (XIGRIS®) ORDER FORM</b>
		<b>***CRITICAL CARE CONSULT REQUIRED FOR USE***</b>
		If you are NOT prescribing drotrecogin alpha please skip to number 24.
		<b>Drotrecogin alfa is available for patients who meet the following criteria:</b>
		a. Systemic Inflammatory Response Syndrome Criteria – Patient must have THREE of the following:
		<input type="checkbox"/> Temperature • 38° C or • 36° C (Pediatrics: • 38.5° C or • 36° C)
		<input type="checkbox"/> Heart rate • 90 beats per minute (Pediatrics: HR • 90 <sup>th</sup> percentile for age)
		<input type="checkbox"/> Respiratory rate • 20 breaths per minute, or PaCO <sub>2</sub> • 32mmHg, or on mechanical ventilation for an acute process (Pediatrics: R • 90 <sup>th</sup> percentile for age or PaCO <sub>2</sub> • 32mmHg)
		<input type="checkbox"/> WBC count • 12,000 or • 4,000/mm <sup>3</sup> , or > 10% immature neutrophils
		b. Organ Dysfunction Criteria – Patient must have at least ONE of the following:
		<input type="checkbox"/> Cardiovascular (Arterial SBP of • 90mmHg or a MAP • 70mmHg for at least 1 hour, despite adequate fluid resuscitation or adequate intravascular volume status or the need for vasopressors to maintain SBP • 90mmHg or MAP • 70mmHg) (Pediatrics: SBP < 10 <sup>th</sup> percentile for age)
		<input type="checkbox"/> Renal (Urine output < 0.5 mL/kg/hour for 1 hour, despite adequate fluid resuscitation) (Pediatrics: Urine output < 1 mL/kg/hour if < 30 kg; Urine output < 30 mL/hour if > 30 kg)
		<input type="checkbox"/> Respiratory (PaO <sub>2</sub> /FiO <sub>2</sub> • 250, or • 200 if the lung was the sole organ meeting the dysfunction criteria)
		<input type="checkbox"/> Hematological (Platelet count of < 80,000/mm <sup>3</sup> or a 50% decrease in the platelet count from the highest value recorded over the previous 3 days)
		<input type="checkbox"/> Unexplained Metabolic Acidosis (pH • 7.3 or base deficit • 5.0 mmol/L and a plasma lactate level > 30 mg/dL)
	18.	
	19.	<b>Relative contraindications to drotrecogin alfa use include:</b>
		• Active internal bleeding, recent (within 3 months) hemorrhagic stroke, recent intracranial or intraspinal surgery, recent severe head trauma, trauma patients with increased risk of life-threatening bleeding, patients with an epidural catheter, patients with intracranial neoplasm, hypersensitivity to drotrecogin.
	20.	<b>Warnings:</b>
		• During Phase III study, incidence of serious bleeding was higher in the treatment group than placebo group; use with caution in patients with an identifiable predisposition to bleeding
	21.	<b>If patient meets the inclusion criteria listed above (within the previous 24 hours), proceed to 'Medication Order'</b>
		<b>Medication Order</b> (check box to order)      Name of critical care attending/fellow authorizing use _____
		<input type="checkbox"/> Drotrecogin alfa 24 mcg/kg/hour continuous IV infusion x 96 hours (maximum dose = 3240 mcg/hour)
	22.	<b>A new order must be written (on Standard Physician Order Form) to continue beyond 96 hours</b>
	23.	<b>Hold infusion for 1 hour before and after any percutaneous procedure; hold for 1 hour before and 12 hours after major surgery</b>
	24.	<b>Reason for not giving drotrecogin alpha:</b>
		<input type="checkbox"/> Active internal bleeding
		<input type="checkbox"/> Hemorrhagic shock
		<input type="checkbox"/> Recent intracranial bleeding or intraspinal surgery
		<input type="checkbox"/> Recent severe head trauma
		<input type="checkbox"/> Patient has an increased risk of life threatening bleeding
		<input type="checkbox"/> Patient has an epidural catheter
		<input type="checkbox"/> Patient has an intracranial neoplasm
		<input type="checkbox"/> Hypersensitivity to drotrecogin alpha
		<input type="checkbox"/> Other (please list reason):

Physician Signature: \_\_\_\_\_ Pager: \_\_\_\_\_

**ACTIVATED PROTEIN C / DROTRECOGIN ALFA (XIGRIS®) ORDER FORM**

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