

# Sepsis Definitions

#### "Sepsis" is defined as a documented or suspected infection with one or more of the following:

#### General variables:

Fever (core temperature >38.3°C)

Hypothermia (core temperature <36°C)

Heart rate >90 min<sup>-1</sup> or >2 SD above the normal value for age

Tachypnea

Altered mental status

Significant edema or positive fluid balance (>20 mL/kg over 24 hrs)

Hyperglycemia (plasma glucose >120 mg/dL) in the absence of diabetes

#### Inflammatory variables:

Leukocytosis (WBC count >12,000 μL<sup>-1</sup>)

Leukopenia (WBC count <4000 μL<sup>-1</sup>)

Normal WBC count with >10% immature forms

Plasma C-reactive protein >2 SD above the normal value

Plasma procalcitonin >2 SD above the normal value

#### Other:

SvO<sub>2</sub>>70%

Cardiac index  $> 3.5 \text{ L} \cdot \text{min}_{-1} \cdot \text{M}_{-23}$ 

# "Severe sepsis" is defined as sepsis associated with organ dysfunction, hypoperfusion or hypotension.

#### Organ dysfunction variables:

Arterial hypoxemia (PaO2/FIO2 <300)

Acute oliguria (urine output <0.5 mL·kg·1·hr·1 or 45 mmol/L for at least 2 hrs)

Creatinine > 2.0 mg/dL

Coagulation abnormalities (INR >1.5 or aPTT >60 secs)

Thrombocytopenia (platelet count <100,000 µL<sup>-1</sup>)

Hyperbilirubinemia (plasma total bilirubin > 2.0 mg/dL or 35 mmol/L)

#### <u>Tissue perfusion variables:</u>

Hyperlactatemia (>2 mmol/L)

#### Hemodynamic variables:

Arterial hypotension (SBP <90 mm Hg, MAP <70, or SBP decrease >40 mm Hg)



#### "Septic shock" is defined as acute circulatory failure unexplained by other causes.

Acute circulatory failure is defined as persistent arterial hypotension (SBP <90 mmHg, MAP <60, or a reduction in SBP >40 mm Hg from baseline despite adequate volume resuscitation).

#### "Bundle" (as in "Sepsis Bundle") is defined as the following:

A group of interventions related to a disease that, when implemented together, result in better outcomes than when implemented individually. The science behind the elements of the bundle is so well-established that their implementation should be considered a generally accepted practice. Bundle components can be easily measured as completed or not completed. As such, the overall bundle — all of the elements taken together — can also be measured as completed or not completed.

# **Evaluation for Severe Sepsis Screening Tool**

	uctions: Use this optional tool wards, or in the ICU.	to scre	en patients for severe	sepsis in the	emergency department,
"* <u>lm</u>	portant: select a date format: _	(d	d/mm/yy) (mm/d	d/yy)	
. Is	the patient's history suggestive	e of a r	new infection?		YesNo
a. b. c.	Pneumonia/ empyema UTI Acute abdominal infection	d. e. f.	Meningitis Skin/soft tissue inflammatic Bone/joint infection	g. on h. i.	
Aı	re any two of following signs &	sympt	oms of infection both p	resent and n	ew to the patient?
a b	Hyperthermia > 38.3 °C (101.0 °F) Hypothermia < 36 °C (96.8 °F)	c. d. e.	Tachycardia > 90 bpm	f. g.	
the	answer is yes to either questio	n 1 or	2, suspicion of infectio	<i>n</i> is present:	<u> </u>
. A	At the physician's discretion obtained any of the following organ diffection that are not considered.  Acutely altered mental status SBP < 90 or MAP < 65 mmHg SpO <sub>2</sub> < 90% on room air or on s Creatinine > 2.0 mg/dl (176.8 mm Bilirubin > 2 mg/dl (34.2 mmol/L) Platelet count < 100,000 Lactate > 2 mmol/L (18.0 mg/dl)	ysfunc I to be upplen mol/L)	ction criteria present at chronic conditions?	a site remote	e from the site of the
	spicion of infection is present A ERE SEPSIS and should be ente				
			Addressograph		
atie	nt Identifier:				
ate ate	of Admission:// format as above).				
me	of Admission:: (24 h	r. cloc	κ)		

### Severe Sepsis Bundles:

### Sepsis Resuscitation Bundle

(To be accomplished as soon as possible and scored over first 6 hours):

- 1. Serum lactate measured.
- 2. Blood cultures obtained prior to antibiotic administration.
- 3. From the time of presentation, broad-spectrum antibiotics administered within 3 hours for ED admissions and 1 hour for non-ED ICU admissions.
- 4. In the event of hypotension and/or lactate > 4 mmol/L (36 mg/dl):
  - a) Deliver an initial minimum of 20 ml/kg of crystalloid (or colloid equivalent\*).
  - b) Apply vasopressors for hypotension not responding to initial fluid resuscitation to maintain mean arterial pressure (MAP)  $\geq$  65 mm Hg.
- 5. In the event of persistent hypotension despite fluid resuscitation (septic shock) and/or lactate > 4 mmol/L (36 mg/dl):
  - a) Achieve central venous pressure (CVP) of  $\geq$  8 mm Hg.
  - b) Achieve central venous oxygen saturation (ScvO<sub>2</sub>) of  $\geq$  70%.\*\*

### Sepsis Management Bundle

(To be accomplished as soon as possible and scored over first 24 hours):

- 1. Low-dose steroids\* administered for septic shock in accordance with a standardized ICU policy.
- 2. Drotrecogin alfa (activated) administered in accordance with a standardized ICU policy.
- Glucose control maintained > lower limit of normal, but < 150 mg/dl (8.3 mmol/L).</li>
- 4. Inspiratory plateau pressures maintained < 30 cm H<sub>2</sub>O for mechanically ventilated patients.

<sup>\*</sup>See the individual chart measurement tool for an equivalency chart.

<sup>\*\*</sup>Achieving a mixed venous oxygen saturation (SvO<sub>2</sub>) of 65% is an acceptable alternative.

### Individual Chart Measurement Tool:

<u>Instructions</u>: Attach this tool to each chart of a patient with severe sepsis, septic shock at the time of data abstraction. This tool can be used for concurrent, prospective, or retrospective data collection. However, individual hospitals are strongly encouraged to choose a single approach and maintain that collection over time. Once all Individual Chart Measurement Tools are gathered for a single month, complete the Monthly Measurement Worksheet to report results.

***	mporta	nt: mark the date format you will be following: (dd/mm/yy) (mm/dd/yy)
1.	answe	nent whether the patient met criteria for severe sepsis or septic shock. Check only one r. Because strict definitions apply it may be helpful to consult the <u>Sepsis Definitions</u> r the <u>Evaluation for Severe Sepsis Screening Tool</u> to ensure accuracy.
		No, does not meet criteria for either severe sepsis or septic shock. Stop data collection. Yes, met criteria for severe sepsis. Continue data collection. Yes, met criteria for septic shock. Continue data collection.
2.	Record	the patient identifier number
3.	individ subsec	on 3 establishes a uniform "time of presentation" for each patient depending upon their lual admission characteristics. The time of presentation will be the basis for answering quent questions and making calculations. Only one statement below (3a, 3b, or 3c) will to a single patient.
		A protocol, protocol form and protocol order set are recommended to facilitate the ent process and the accurate recording of timelines.
	За.	For patients admitted to the ICU from the ED meeting criteria for severe sepsis or septic shock, record the time of triage in the emergency department as the time of presentation.
		Not applicable. Proceed to 3b. Applicable, record time of presentation below and proceed to question 4.
	3h	For nationts transferred to the ICII from units other than the ED:

- 3b. For patients transferred to the ICU from units other than the ED:
  - Preferred: if the resuscitation and management of severe sepsis was annotated as beginning on the transferring unit, record the time and date of that annotation as the time of presentation.
  - Default: if the resuscitation and management of severe sepsis was not in annotated as beginning on the transferring unit, record the ICU admission date and time as the time of presentation.

<u>Note</u>: it is critical to establish whether there was reasonable and straightforward annotation of the time of initiation of efforts to manage severe sepsis on the ward prior to ICU transfer. Otherwise, no credit can be assigned for key interventions performed prior to the default time of presentation, the time of ICU admission. Annotation may include a practitioner's note, a practitioner's timed and dated orders, a nurse's timed and dated records documenting discussion of severe sepsis with a practitioner, timed records initiating referral to the ICU for severe sepsis.

		Not applicable. Proceed to question 3c.
		Applicable; the annotated time and date for the resuscitation and management of sepsis on the transferring unit is recorded below as the time of presentation.  Proceed to question 4.
		Applicable; the ICU admission date and time is recorded below as the time of presentation. Proceed to question 4.
		3c. For patients admitted to the ICU with a diagnosis other than sepsis and who subsequently develop severe sepsis or septic shock on the same ICU stay, record the annotated time and date of the beginning of the resuscitation and management of severe sepsis as the time of presentation.
		Not applicable. Stop data collection, time of presentation cannot be accurately determined. If data is being collected concurrently or prospectively, the patient may remain on the sepsis protocol without further data collection.
		Applicable, record time of presentation below and proceed to question 4.
***	Time	of Presentation: / / (date format as above) : (24 hour clock).***
		, , , , , , , , , , , , , , , , , , , ,
4.	Doc	cument whether serum lactate was obtained:
		No. Proceed to question 5.
		No. Proceed to question 5 Yes. Place a mark in Box 1 on line 16 of this document. Proceed to question 4a.
		Record the value serum lactate value if obtained: mmol/L or mg/dl
	4b.	Record date and time of serum lactate collection:
		// (date format as above): (24 hour clock).
5.	Doo	cument whether the patient received a broad-spectrum antibiotic:
		_ No. Proceed to question 7.
		Yes. Proceed to question 5a.
		Name of Antibiotic(s):
	5b.	Date and time of first broad-spectrum antibiotic administration:
	_	/(date format as above):(24 hour clock).
		Calculate the difference between line 3, time of presentation above, and line 5b in hours and minutes: Difference: hours minutes
		Multiply the HOURS ONLY on line 5c above x 60
		Time in minutes to broad spectrum antibiotic administration for this patient: add the total of line 5d above to the number of MINUTES ONLY listed on line 5c above:
	5f.	If item 3a above is marked applicable, was the number of minutes on line 5e above ≤ 180
		minutes:
		No. Proceed to question 6.
	<b>.</b>	Yes. Place a mark in Box 3 on line 16 of this document. Proceed to question 6.
	_	If item 3b or 3c is marked applicable, was the number of minutes on line 5e above ≤ 60 minutes:
		No. Proceed to question 6.
		Yes. Place a mark in Box 3 on line 16 of this document. Proceed to question 6.

6.	Document date and time of blood culture collection.
	If not collected, enter "No" on 6a and proceed to question 7 / / (date format as above) : (24 hour clock). Proceed to question 6a.
	6a. Document whether the time and date listed on 6 above is earlier than the time and date listed on line 5b above:
	No. Proceed to question 7.
	Yes. Place a mark in Box 2 on line 16 of this document. Proceed to question 7.
7.	Answer the following questions regarding resuscitation of severe sepsis or septic shock:
	7a. Document whether the patient was hypotensive <u>and/or</u> if serum lactate was > 4 mmol/L (30 mg/dl) on line 4a of this document:
	No. Place a mark in Box 4, 5, 6, 7 on line 16 of this document. Place a mark in Box A on line 17 of this document. Proceed to question 11.
	Yes. Proceed to question 7b.
	7b. Document the basis for the diagnosis of hypotension, if present:
	SBP < 90 mm Hg
	MAP < 65 mm Hg Note: $MAP = (2 \times diastolic \ pressure + systolic \ pressure) / 3$
	SBP decrease of ≥ 40 mm Hg from known baseline
	7c. Document whether initially the patient received ≥ 20 ml/kg of crystalloid or ≥ an equivalent amount of colloid in response to hypotension or lactate > 4 mmol/L (36 mg/dl):

Cyrstalloid/Colloid Equiva	lency Chart: <sup>1</sup>
Normal Saline	20 ml/kg
Lactated Ringer's Solution	20 ml/kg
Albumin	0.24 grams/kg
4-5% Albumin	5.2 ml/kg
20-25% Albumin	1.1 ml/kg
Hetastarch	0.29 grams/kg
3% Hetastarch	9.7 ml/kg
6% Hetastarch	4.8 ml/kg
10% Hetastarch	2.9 ml/kg
Pentastarch	0.30 grams/kg
10% Pentastarch	3 ml/kg
10% Dextran-40	0.30 grams/kg (3ml/kg)
3% Dextran-60, 6% Dextran-70	0.19 grams/kg
3% Dextran-60	6.3 ml/kg
6% Dextran-70	3.1 ml/kg
Gelatins (succinylated & crosslinked 2.5, 3.0, 4.0%; urea-linked 3.5%)	0.23 grams/kg

Adapted from: Evidence-based Colloid Use in the Critically III: American Thoracic Society Consensus Statement. Am J Respir Crit Care Med. 2004. Vol 170:1247-1259. For percentage solutions, listed ml/kg are calculated from the g/kg data.

No. Record "No" on lines 7f, 8b, 9b and 10 below. Proceed to question 11.

	Yes. Place a mark in Box 4 on line 16 of this document. Proceed to question 7d.
	Document whether MAP remained ≥ 65 in response to the initial fluid resuscitation described in 7c:
	<ul> <li>ii No. Proceed to question 7e.</li> <li>iii Yes, if lactate was ≤ 4 mmol/L (36 mg/dl) on line 4a of this document place a mark in Box 5, Box 6 and Box 7 on line 16 of this document. Proceed to question 10.</li> <li>iii Yes, if lactate was &gt; 4 mmol/L (36 mg/dl) on line 4a of this document, proceed to question 8.</li> </ul>
7e.	Document whether the patient received vasopressors:  No. Record "No" on lines 7f, 8b, 9b and 10 below. Proceed to question 11.  Yes. Place a mark in Box 5 on line 16 of this document. Proceed to question 7f.
7f.	Document whether the MAP remained ≥ 65 mm Hg without the use of vasopressors:
	Note: If no evidence for removal of vasopressors can be found, mark item 7f "no" and proceed to question 8.
	<ul> <li>ii No. Proceed to question 8.</li> <li>iii Yes, if lactate was ≤ 4 mmol/L (36 mg/dl) on line 4a of this document place a mark in Box 6 and Box 7 on line 16 of this document. Proceed to question 10.</li> <li>iii Yes, if lactate was &gt; 4 mmol/L (36 mg/dl) on line 4a of this document, proceed to question 8.</li> </ul>
Doc	<ul> <li>cument date and time CVP first ≥ 8 mm Hg within 24 hours:</li> <li>CVP not obtained or never ≥ 8 mm Hg within 24 hours. Record line 8b as "No" and proceed to question 9.</li> </ul>
	e://
	Calculate the difference between line 3, time of presentation, and line 8 above in hours and minutes: Difference:: (hours:minutes).
	Document whether line 8a is ≤ 6 hours.  No. Proceed to question 9.  Yes. Place a mark in Box 6 on line 16 of this document. Proceed to question 9.
	sument date and time $ScvO_2$ first $\geq 70\%$ (or $SvO_2 \geq 65\%$ ) within 24 hours: $\_ScvO_2$ not obtained or never $\geq 70\%$ (or $SvO_2 \geq 65\%$ ) within 24 hours. Record line 9b as "No" and proceed to question 10.
	e:/ (date format as above) <b>Time</b> : : (24 hour clock). ceed to question 9a.
	Calculate the difference between line 3, time of presentation, and line 9 above in hours and minutes: Difference:: (hours:minutes).
9b.	Document whether line 9a is ≤ 6 hours.  No. Proceed to question 10.

8.

9.

	not have persistent arterial hy to question 11.	ot applicable because the patient's MAP was ≥ 65 potension. Place a mark in Box A on line 17 and	
	No. Proceed to question 10b.		
10b.		dardized ICU policy regarding low-dose stero	oid
	administration for septic shock: No. Proceed to question 11.		
	Yes. Proceed to question 10c		
		esent. Place a mark in Box A on line 17 below.	Proce
	question 11.		
	question 11.		
10d.	Document whether low-dose stero	oids were administered:	
10d.	Document whether low-dose stero		
10d.	Document whether low-dose stero	oids were administered:  daily dose of 200–300 mg of hydrocortisone	or
10d.	Document whether low-dose stero		or
10d.	Document whether low-dose steroids refer to a equivalent.		or
10d.	Document whether low-dose steroids refer to a equivalent.	daily dose of 200–300 mg of hydrocortisone	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid	daily dose of 200–300 mg of hydrocortisone	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid:	daily dose of 200–300 mg of hydrocortisone  Equivalency Chart:  Equivalent TOTAL DAILY dose:	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid Steroid: Hydrocortisone	daily dose of 200–300 mg of hydrocortisone  Equivalency Chart:  Equivalent TOTAL DAILY dose:  200 – 300 mg	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid Steroid: Hydrocortisone Dexamethasone	daily dose of 200–300 mg of hydrocortisone  Equivalency Chart:  Equivalent TOTAL DAILY dose:  200 – 300 mg  8 – 12 mg	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid Steroid: Hydrocortisone Dexamethasone Prednisone	daily dose of 200–300 mg of hydrocortisone  Equivalency Chart:  Equivalent TOTAL DAILY dose:  200 – 300 mg  8 – 12 mg  50 – 75 mg	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid Steroid: Hydrocortisone Dexamethasone Prednisolone Prednisolone	daily dose of 200–300 mg of hydrocortisone  Equivalency Chart:  Equivalent TOTAL DAILY dose:  200 – 300 mg  8 – 12 mg  50 – 75 mg  50 – 75 mg	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisolone Methylprednisolone	daily dose of 200–300 mg of hydrocortisone delivations and the second se	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid Steroid: Hydrocortisone Dexamethasone Prednisone Prednisolone Methylprednisolone Cortisone	daily dose of 200–300 mg of hydrocortisone descriptions of 200–300 mg of hydrocortisone descriptions of 200–300 mg  8 – 12 mg  50 – 75 mg  40 – 60 mg  250 – 375 mg	or
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisone Prednisolone Methylprednisolone Cortisone Triamcinolone Betamethasone	daily dose of 200–300 mg of hydrocortisone           Equivalency Chart: <sup>2</sup> Equivalent TOTAL DAILY dose:           200 – 300 mg           8 – 12 mg           50 – 75 mg           40 – 60 mg           250 – 375 mg           40 – 60 mg           6 – 10 mg	
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisone Prednisolone Methylprednisolone Cortisone Triamcinolone Betamethasone	daily dose of 200–300 mg of hydrocortisone           Equivalency Chart: <sup>2</sup> Equivalent TOTAL DAILY dose:           200 – 300 mg           8 – 12 mg           50 – 75 mg           40 – 60 mg           250 – 375 mg           40 – 60 mg           40 – 60 mg	
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisolone Methylprednisolone Cortisone Triamcinolone Betamethasone	daily dose of 200–300 mg of hydrocortisone           Equivalency Chart: <sup>2</sup> Equivalent TOTAL DAILY dose:           200 – 300 mg           8 – 12 mg           50 – 75 mg           40 – 60 mg           250 – 375 mg           40 – 60 mg           6 – 10 mg	
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisone Prednisolone Methylprednisolone Cortisone Triamcinolone Betamethasone	Equivalency Chart: <sup>2</sup> Equivalent TOTAL DAILY dose:  200 – 300 mg  8 – 12 mg  50 – 75 mg  40 – 60 mg  250 – 375 mg  40 – 60 mg  40 – 60 mg  6 – 10 mg  andbook of Clinical Drug Data, 6th ed. Drug Intelligence Pub,	
10d.	Note: low-dose steroids refer to a equivalent.  Steroid: Hydrocortisone Dexamethasone Prednisone Prednisolone Methylprednisolone Cortisone Triamcinolone Betamethasone  2 Adapted from: Knoben JE, Anderson PO. Ha	Equivalency Chart: <sup>2</sup> Equivalent TOTAL DAILY dose:  200 – 300 mg  8 – 12 mg  50 – 75 mg  40 – 60 mg  250 – 375 mg  40 – 60 mg  40 – 60 mg  6 – 10 mg  andbook of Clinical Drug Data, 6th ed. Drug Intelligence Pub,	

	10f.	Document whether the time and date on 10d is $\leq$ 24 hours from the time of presentation listed on item 10e.
		No. Proceed to question 11.
		Yes. Place a mark in Box A on line 17 and proceed to question 11.
11.	Ansv	wer the following questions regarding Drotrecogin alfa (activated) administration:
	11a.	Document whether there is a standardized ICU policy regarding Drotrecogin alfa (activated) administration:
		No. Proceed to question 12.
		Yes. Proceed to question 11b.
	11b.	Indicate whether there is documentation that the patient <u>did not merit</u> Drotrecogin alfa (activated) administration based upon the standardized protocol:
		No documentation is present. Proceed to question 11c.
		Yes there is documentation present. Place a mark in Box B on line 17 below. Proceed to question 12.
	11c.	Document whether Drotrecogin alfa (activated) was administered: No. Proceed to question 12.
		Yes. Record date and time below. Proceed to question 11d.
		//(date format as above) : (24 hour clock)
		(
	11d.	Time of presentation: from line 3 above:
		/(date format as above) : (24 hour clock)
	11e.	Document whether the time and date on 11c is $\leq$ 24 hours from the time of presentation listed on item 11d.
		No. Proceed to question 12.
		Yes. Place a mark in Box B on line 17 and proceed to question 12.
12.	Docu	ument the median glucose* value within 24 hours of the time of presentation:
	Medi	ian glucose: mg/dl or mmol/L
		d only if median glucose is < 150 mg/dl (8.3 mmol/L) place a mark in Box C on line 17 of this ment. Proceed to question 12a.
		Document the lower limit of normal for serum glucose at your institution:  Document the total number of measurements that fell below the lower limit of normal within 24 hours from the time of presentation for this patient:
	* Ref	fer to the optional <i>Median Glucose Tool,</i> if necessary.
13.		ument the median inspiratory plateau pressure (IPP)* achieved within 24 hours of time of entation:
		Not applicable because the patient was not mechanically ventilated. Place a mark in Box D on line 17 of this document. Proceed to question 14.
	Medi	ian IPP: If and only if < 30 cm H <sub>2</sub> 0, place a mark in Box D on line 17 of this document.

14.	Date and	time of he	ospital dis	scharge:			
	/_	/	(date form	at as abov	re):	(24	hour clock)
15.	Status at	hospital	discharge	:Aliv	/e Dec	ceased	
16.	Boxes 1 t	through 7	:				
	Box 1	Box 2	Box 3	Box 4	Box 5	Box 6	Box 7
17.	Boxes A	through [	):				
	Box A	Box B	Box C	Box D			

\* Refer to the optional *Median IPP Calculation Tool*, if necessary.

## Monthly Measurement Worksheet:

<u>Instructions:</u> Gather the individual chart measurement tools that have been collected in a particular month and complete the following form.

\*\*Note: Where blanks are found on the Individual Chart Measurement Tool as you tally results, this is by design. Blank items should not factor into your calculations.

1)		al number of patients selected for monthly reporting. Each patient should have an individual chart asurement tool completed:
2)	Rel	iability of compliance with the sepsis resuscitation bundle element 1: lactate measured:
		Tally the number of affirmative answers to question 4 on the individual chart measurement tool: Reliability of compliance with the sepsis resuscitation bundle element 1: divide line 2a from this worksheet by line 1 from this worksheet:
3)	Qu	ality Indicator #1, blood cultures collected before broad-spectrum antibiotic administration:
		Tally the number of affirmative answers to question 6a on the individual chart measurement tool:  Reliability of compliance with sepsis resuscitation bundle element 2, blood cultures before antibiotics administered: divide line 3a from this worksheet by line 1 from this worksheet:*
	Nu Dei	merator for Improvement Tracker (from line 3a on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):
		ne reliability of compliance with the second bundle element is the same as the final value for quality cator #1 in this instance.
4)		ality Indicator #2, median time in minutes to broad-spectrum antibiotic(s) administration from time of sentation:
	b) c)	Tally the number of affirmative answers to question 5f on the individual chart measurement tool: Tally the number of affirmative answers to question 5g on the individual chart measurement tool: Reliability of compliance with sepsis resuscitation bundle element 3, from the time of presentation, broad-spectrum antibiotics administered within 3 hours for ED admissions and 1 hour for non-ED ICU admissions: add lines 4a and 4b on this worksheet and divide the total by line 1 on this worksheet: Tally the median number of minutes to antibiotic administration as follows:
		Arrange each value listed on line 5e of the individual chart measurement tools for each patient in numerical sequence from the smallest to the largest:
		Smallest
		Largest
		The median is the middle value of the set if the number of measurements is odd and the average of the

Median number of minutes to broad spectrum antibiotic administration. Report in Improvement

central pair if the number is even.

Tracker: \_\_\_\_

5)		liability of compliance with sepsis resuscitation bundle element 4a: for hypotension and/or lactate inmol/L (36 mg/dl), deliver an initial minimum of 20 ml/kg of crystalloid (or colloid equivalent):
	a) b)	Tally the number of charts with an affirmative answer to question 7a on the individual chart measurement tool:  Tally the number of charts with affirmative answers to question 7c on the individual chart measurement
	c)	tool Reliability of compliance with sepsis resuscitation bundle element 4a: divide line 5b on this worksheet by line 5a on this worksheet
6)	hyp	liability of compliance with sepsis resuscitation bundle element 4b: apply vasopressors for potension not responding to initial fluid resuscitation to maintain mean arterial pressure (MAP) $\geq$ 65 and $\geq$ 65 and $\geq$ 65 and $\geq$ 65 are the complex constants.
	a)	Tally the number of charts with negative answers to question 7d on the individual chart measurement tool:
	b)	Tally the number of charts with affirmative answers to question 7e on the individual chart measurement tool:
	c)	Reliability of compliance with sepsis resuscitation bundle element 4b: divide line 6b on this worksheet by line 6a on this worksheet:
7)		<b>ality Indicator #3</b> , following septic shock or lactate > 4 mmol/L (36 mg/dl) identification, central venous ssure (CVP) of $\geq$ 8 mm Hg achieved within 6 hours of presentation time:
	<ul><li>a)</li><li>b)</li><li>c)</li><li>d)</li><li>e)</li></ul>	Number of patients in septic shock: tally the number of charts marked with a negative response to question 7f on the individual chart measurement tool
		merator for Improvement Tracker (from line 7d on this worksheet): nominator for Improvement Tracker (from line 7c on this worksheet):
		he reliability of compliance with sepsis resuscitation bundle element 5a is the same as the final value for ality indicator #3 in this instance.
8)		<b>ality Indicator #4,</b> following septic shock or lactate > 4 mmol/L (36 mg/dl) identification, $ScvO_2 \ge 70\%$ $SvO_2 \ge 65\%$ ) achieved within 6 hours of presentation time:
	a) b) c)	Total from line 7c on this worksheet:Number of patients who are in septic shock or who have lactate greater than 4 mmol/L (36 mg/dl) for whom $ScvO_2$ is $\geq 70\%$ (or $SvO_2$ is $\geq 65\%$ ) within 6 hours of presentation: tally the number of charts with an affirmative answer to question 9b on the individual chart measurement tool: Reliability of compliance with sepsis resuscitation bundle element 5b, in the event of persistent hypotension despite fluid resuscitation (septic shock) and/or lactate > 4 mmol/L (36 mg/dl), achieve central venous oxygen saturation ( $ScvO_2$ ) of 70% (or mixed venous oxygen saturation ( $SvO_2$ ) of 65%): divide line 8b on this worksheet by line 8a on this worksheet:*

		merator for Improvement Tracker (from line 8b on this worksheet): nominator for Improvement Tracker (from line 8a on this worksheet):
		ne reliability of compliance with sepsis resuscitation bundle element 5b is the same as the final value for ality indicator #4 in this instance.
9)		ality Indicator #5, low-dose steroids administered in accordance with a standardized ICU policy for otic shock over the first 24 hours following the time of presentation:
	b)	Total number of patients with septic shock: line 7a on this worksheet: Tally the number of affirmative answers to question 10c on the individual chart measurement tool: Tally the number of affirmative answers to question 10f on the individual chart measurement tool: Number of patients for whom steroids were administered in accordance with a standardized ICU policy within 24 hours of presentation: Add lines 9b on this worksheet and 9c on this worksheet: Reliability of compliance with sepsis management bundle element 1, low-dose steroids administered for septic shock in accordance with a standardized ICU policy: Divide line 9d on this worksheet by line 9a on this worksheet:*
	Nu De	merator for Improvement Tracker (from line 9d on this worksheet): nominator for Improvement Tracker (from line 9a on this worksheet):
		ne reliability of compliance with sepsis management bundle element 1 is the same as the final value for ality indicator #5 in this instance.
10)		ality Indicator #6, Drotrecogin alfa (activated) administered in accordance with a standardized ICU icy for septic shock over the first 24 hours following time of presentation:
		Tally the number of affirmative answers to question 11b on the individual chart measurement tool:  Tally the number of affirmative answers to question 11e on the individual chart measurement tool:  Number of patients for whom Drotrecogin alfa (activated) was administered in accordance with a standardized ICU policy within 24 hours of presentation: Add lines 10a on this worksheet and 10b on this worksheet:  Reliability of compliance with sepsis management bundle element 2, Drotrecogin alfa (activated) administered in accordance with a standardized ICU policy: divide line 10c on this worksheet by line 1 on this worksheet:  **  **  **  **  **  **  **  **  **
		merator for Improvement Tracker (from line 10c on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):
		ne reliability of compliance with sepsis management bundle element 2 is the same as the final value for ality indicator #6 in this instance.
11)		ality Indicator #7, glucose control maintained $\geq$ the lower limit of normal and with median value < 150 /dl (8.3 mmol/L) over the first 24 hours following presentation with severe sepsis, septic shock:
	a)	Tally the number of patients for whom median glucose was recorded to be < 150 on line 12 of the individual chart measurement tool <u>AND</u> for whom no episodes of hypoglycemia were recorded on line 12b:
	b)	Reliability of compliance with sepsis management bundle element 3, glucose control maintained > lower limit of normal, but < 150 mg/dl (8.3 mmol/L): divide line 11a on this worksheet by line 1 on this worksheet:
	Nu De	merator for Improvement Tracker (from line 11a on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):

		ne reliability of compliance with sepsis management bundle element 3 is the same as the final value for ality indicator #7 in this instance.
12)		ality Indicator #8, median inspiratory plateau pressure (IPP) less than 30 cm $H_2O$ over the first 24 urs in patients presenting with severe sepsis, septic shock who require mechanical ventilation:
	b)	Total from line 1 on this worksheet: Tally the number of cases marked on line 13 of the individual chart measurement tool as not applicable because the patient was not mechanically ventilated: Number of mechanically ventilated patients: subtract line 12b on this worksheet from line 12a on this worksheet: Tally the number of patients for whom median IPP was maintained less than 30 cm $H_20$ on line 13 of the individual chart measurement tool: Reliability of compliance with sepsis management bundle element 4, for mechanically ventilated patients, plateau pressures maintained < 30 cm $H_20$ , divide line 12d by line 12c: *
		merator for Improvement Tracker (from line 12d on this worksheet):nominator for Improvement Tracker (from line 12c on this worksheet):
		ne reliability of compliance with sepsis management bundle element 4 is the same as the final value for ality indicator #8 in this instance.
13)	Qu	ality Indicator #9, reliability of compliance with ALL elements of the severe sepsis resuscitation bundle:
	,	Tally the number of charts with ALL boxes (1 through 7) in question 16 on the individual chart measurement tool marked:  Aggregate reliability for the sepsis resuscitation bundle: divide line 13(a) by line 1
		merator for Improvement Tracker (from line 13a on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):
14)		ality Indicator #10, reliability of compliance with ALL elements of the severe sepsis management adle:
	,	Tally the number of charts with ALL boxes (A through D) in question 17 on the individual chart measurement tool marked:  Aggregate reliability of sepsis management bundle: divide line 14(a) by line 1
		merator for Improvement Tracker (from line 14a on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):
15)	Qu	ality Indicator #11, overall mortality due to severe sepsis, septic shock:
	a)	Tally the number of patients marked as deceased from line 15 of the individual chart measurement tool:
	b)	Overall mortality due to severe sepsis, septic shock: divide line 15(a) by line 1.
		merator for Improvement Tracker (from line 15a on this worksheet): nominator for Improvement Tracker (from line 1 on this worksheet):

## Median Glucose Calculation Tool:

Use this optional tool to assist in determining the median glucose level within the first 24 hours from time of presentation.

cord the patie	nt id	entif	ier n	umb	er			-								
Worksheet:																
List all glucos	e val	ues o	obtaiı	ned ir	ո 24	hours	S:									
Time (24° clock)																
Glucose																
Time (24° clock)																
Glucose																
To obtain the smallest to th			lucos	se, fir	st an	range	the	value	es in	nume	eric	al sec	quen	ce fr	om t	he
Smallest																Larges
The median is the middle value of the set if the number of measurements is odd and the average of the central pair if the number is even.																
edian glucose:	l	m(	g/dl d	or	m	ımol/	L									
cord results o	n the	Sev	ere	Seps	is Q	ualit	v Me	asur	eme	nt To	ol ı	ınde	r Me	dian	Glu	cose.

## Median Plateau Pressure Tool:

Use this optional tool to assist in determining the median inspiratory plateau pressure for mechanically ventilated patients with severe sepsis within the first 24 hours from time of presentation.

Worksheet:																
List all IPP va	lues o	btair	ned ir	1 24 h	nours											
Time			1						1		- 1					
(24° clock)																
IPP																
					ı		- I									
Time																
(24° clock)																
IPP																
	media	n IP	P, firs	st arra	ange	the v	alues	s in n	ume	rical s	equ	ience	fron	n the s	smal	lest to
To obtain the the largest:  Smallest																Large
the largest:				ue of	the s	set if t	he nı	umbe	er of	meas	urer	ments	is o	odd and	d the	
the largest: Smallest	the m	niddl	 e val					umbe	er of	meas	urer	ments	is o	dd an	d the	
the largest: Smallest The median is	the m	niddl	 e val					umbe	er of	measi	urer	ments	is o	dd an	d the	

## Severe Sepsis Quality Indicators

Quality Indicator #1	Definition of Indicator	Specifications
Blood cultures	The percent of patients	Numerator: number of patients who had
collected before	presenting with severe	blood cultures collected before broad-
broad-spectrum	sepsis, septic shock who	spectrum antibiotic administration following
antibiotic	had blood cultures	presentation with severe sepsis, septic shock
administration	collected prior to broad-	
	spectrum antibiotic	Denominator: number of patients presenting
	administration	with severe sepsis, septic shock

Quality Indicator #2	Definition of Indicator	Specifications
Median time in minutes to broad-spectrum antibiotic(s) administration from	Median time to administration of broad-spectrum antibiotic(s).	For patients presenting with severe sepsis, septic shock, median time in minutes to broad-spectrum antibiotic(s) administration from time of presentation.
time of presentation.		For this indicator, time of presentation is determined as follows:

- (i) If the patient presented to the ED with severe sepsis, septic shock, the time of presentation is ED triage time.
- (ii) If the management of severe sepsis, septic shock was annotated as beginning on a unit transferring the patient to the ICU, the annotated time and date of the initiation of the resuscitation is the time of presentation.
- (iii) If the management of severe sepsis, septic shock was NOT annotated as beginning on a unit transferring the patient to the ICU, the recorded ICU admission time is the default value for time of presentation.
- (iv) If the patient is newly treated for severe sepsis, septic shock while in the ICU greater than 24 hours after admission to the ICU, the annotated time and date of the initiation of resuscitation for the management of severe sepsis, septic shock is the time of presentation.

Quality Indicator #3	Definition of Indicator	Specifications
Following septic	The percent of patients	Numerator: the number of patients with
shock or lactate > 4	for whom a CVP goal of >	severe sepsis with lactate > 4 mmol/L (36
mmol/L (36 mg/dl)	8 mm Hg was achieved	mg/dl) or septic shock for whom a CVP goal
identification, central	within 6 hours following	≥ 8 mm Hg was achieved within 6 hours of
venous pressure	septic shock or lactate >	presentation time.
(CVP) of <u>&gt;</u> 8 mm Hg	4 mmol/L (36 mg/dl)	
achieved within 6	identification.	Denominator: the number of patients with
hours of		severe sepsis with lactate > 4 mmol/L (36
presentation time.		mg/dl) or septic shock.
		Evaluaian; nationto with non acyora consis or
		Exclusion: patients with non-severe sepsis or severe sepsis with lactate < 4 mmol/L (36
		mg/dl).
		mg/ui).

Quality Indicator #4	Definition of Indicator	Specifications
Following septic	The percent of patients	Numerator: the number of patients with
shock or lactate > 4	following septic shock or	severe sepsis with lactate > 4 mmol/L (36
mmol/L (36 mg/dl)	lactate > 4 mmol/L (36	mg/dl) or septic shock for whom a ScVO <sub>2</sub>
identification, central	mg/dl) identification for	goal $\geq$ 70% (or SvO <sub>2</sub> goal $\geq$ 65%) is achieved
venous oxygen	whom the $ScvO_2$ is $\geq 70\%$	within 6 hours of presentation time.
saturation (ScvO <sub>2</sub> )* $\geq$	(or $SvO_2$ is $\geq$ 65%) within	
70% achieved within	6 hours of presentation	Denominator: the number of patients with
6 hours of	time.	severe sepsis with lactate > 4 mmol/L (36
presentation time.		mg/dl) or septic shock.
*mixed venous		Fuelveien estimate with more excessions
oxygen saturation		Exclusion: patients with non-severe sepsis or
$(SvO_2) \ge 65\%$ may		severe sepsis with lactate < 4 mmol/L (36
be substituted.		mg/dl).

Quality Indicator #5	Definition of Indicator	Specifications
Low-dose* steroids	The percent of patients	Numerator: number of patients with septic
administered in	with septic shock for	shock for whom low-dose steroids were
accordance with a	whom low-dose steroids	administered in accordance with a
standardized ICU	were administered in	standardized ICU policy within 24 hours
policy for septic	accordance with a	following time of presentation.
shock over the first	standardized ICU policy	
24 hours following	within 24 hours following	Denominator: total number of patients with
time of presentation.	the time of presentation.	septic shock.
		*Low-dose steroids refer to a daily dose of
		200–300 mg of hydrocortisone or equivalent.

Quality Indicator #6	Definition of Indicator	Specifications
Drotrecogin alfa	The percent of patients	Numerator: number of patients with severe
(activated)	with severe sepsis or	sepsis or septic shock for whom Drotrecogin
administered for	septic shock for whom	alpha (activated) was administered in
septic shock in	Drotrecogin alpha	accordance with a standardized ICU policy
accordance with a	(activated) was	within 24 hours following the time of
standardized ICU	administered in	presentation.
policy over the first	accordance with a	
24 hours following	standardized ICU policy	Denominator: total number of patients
time of presentation.	within 24 hours following	presenting with severe sepsis, septic shock
	the time of presentation.	
		Exclusion: non-severe sepsis

Quality Indicator #7	Definition of Indicator	Specifications

Glucose control maintained greater than the lower limit of normal and with median value < 150	The percent of patients with severe sepsis, septic shock for whom serum glucose is maintained greater than the lower	Numerator: number of patients with serum glucose maintained greater than the lower limit of normal and with a median value < 150 mg/dl (8.3 mmol/L) over the first 24 hours following presentation with severe
mg/dl (8.3 mmol/L) over the first 24 hours following presentation with severe sepsis, septic shock.	limit of normal and with a median value < 150 mg/dl (8.3 mmol/L) over the first 24 hours.	sepsis, septic shock  Denominator: number of patients presenting with severe sepsis, septic shock

Quality Indicator #8	Definition of Indicator	Specifications
Median inspiratory plateau pressure (IPP) less than 30 cm H <sub>2</sub> O over the first the first 24 hours in patients presenting with severe	The percent of patients requiring mechanical ventilation who have a median IPP < 30 cm H <sub>2</sub> O over the first 24 hours following presentation with	Numerator: number of mechanically ventilated patients with severe sepsis, septic shock who had a median IPP < 30 cm H₂O over the first 24 hours following presentation with severe sepsis, septic shock.
sepsis, septic shock that require mechanical ventilation.	severe sepsis, septic shock	Denominator: number of mechanically ventilated patients presenting with severe sepsis, septic shock
		Exclusion: Patients not mechanically ventilated.

Definition of Measure	Specifications
The percent of cases of	Numerator: the number of cases of severe sepsis, septic shock that completed all
shock that completed	applicable severe sepsis resuscitation bundle
all applicable severe	elements.
•	Denominator: total number of patients with
	severe sepsis, septic shock
	Exclusion: non-severe sepsis
	The percent of cases of severe sepsis, septic shock that completed

Quality Indicator #10	Definition of Measure	Specifications
Reliability of compliance with all elements of the severe	The percent of cases of severe sepsis, septic shock that completed	Numerator: the number of cases of severe sepsis, septic shock that completed all applicable severe sepsis management
sepsis management bundle.	all applicable severe sepsis management	bundle elements.
	bundle elements	Denominator: total number of patients with severe sepsis, septic shock
		Exclusion: non-severe sepsis

Quality Indicator #11	Definition of Measure	Specifications
Mortality due to severe sepsis, septic shock	The percent of cases of death due to severe sepsis, septic shock	Numerator: the number of cases of death due to severe sepsis, septic shock
	,	Denominator: total number of patients with severe sepsis, septic shock
		Exclusion: non-severe sepsis