


GUIDELINES FOR MANAGEMENT OF SEVERE SEPSIS AND SEPTIC SHOCK

 **Surviving Sepsis Campaign** This is a summary of the Surviving Sepsis Campaign International Guidelines for Management of Severe Sepsis and Septic Shock: 2008, condensed from Dellinger RP, Levy MM, Carlet JM, et al: Surviving Sepsis Campaign: International guidelines for management of severe sepsis and septic shock. *Intensive Care Medicine* (2008) 34:17-60 and *Crit Care Med* 2008; 36(1) 296-327. This version does not contain the rationale or appendices contained in the primary publication. The SSC guidelines do not cover every aspect of managing critically ill patients, and their application should be supplemented by generic best practice and specific treatment as required. Please refer to the guidelines for additional information at www.survivingsepsis.org

Strength of recommendation and quality of evidence have been assessed using the GRADE criteria, presented in brackets after each guideline.

For added clarity:

- ◆ Indicates a strong recommendation or “we recommend”
- ◇ Indicates a weak recommendation or “we suggest”

SSC GUIDELINES HAVE BEEN ENDORSED BY

American Association of Critical-Care Nurses
 American College of Chest Physicians
 American College of Emergency Physicians
 Canadian Critical Care Society
 European Society of Clinical Microbiology and Infectious Diseases
 European Society of Intensive Care Medicine
 European Respiratory Society
 Indian Society of Critical Care Medicine
 International Sepsis Forum
 Japanese Association for Acute Medicine
 Japanese Society of Intensive Care Medicine
 Society of Critical Care Medicine
 Society of Hospital Medicine
 Surgical Infection Society
 World Federation of Critical Care Nurses
 World Federation of Societies of Intensive and Critical Care Medicine.
 Participation and endorsement by German Sepsis Society
 and Latin American Sepsis Institute.

The Surviving Sepsis Campaign is a collaboration of the European Society of Intensive Care Medicine, the International Sepsis Forum, and the Society of Critical Care Medicine.



Society of
Critical Care Medicine
The Intensive Care Professionals



January 2008

Initial resuscitation (first 6 hours)

- ◆ Begin resuscitation immediately in patients with hypotension or elevated serum lactate ≥ 4 mmol/L; do not delay pending ICU admission. (1C)
 - ◆ Resuscitation goals: (1C)
 - Central venous pressure (CVP) 8–12 mm Hg*
 - Mean arterial pressure ≥ 65 mm Hg
 - Urine output ≥ 0.5 mL.kg⁻¹.hr⁻¹
 - Central venous (superior vena cava) oxygen saturation $\geq 70\%$, or mixed venous $\geq 65\%$
 - ◇ If venous O₂ saturation target not achieved: (2C)
 - consider further fluid
 - transfuse packed red blood cells if required to hematocrit of $\geq 30\%$ and/or
 - dobutamine infusion max 20 μ g.kg⁻¹.min⁻¹
- * A higher target CVP of 12-15 mmHg is recommended in the presence of mechanical ventilation or pre-existing decreased ventricular compliance.

Diagnosis

- ◆ Obtain appropriate cultures before starting antibiotics provided this does not significantly delay antimicrobial administration. (1C)
 - Obtain two or more blood cultures (BCs)
 - One or more BCs should be percutaneous
 - One BC from each vascular access device in place ≥ 48 hours
 - Culture other sites as clinically indicated
- ◆ Perform imaging studies promptly in order to confirm and sample any source of infection if safe to do so. (1C)

Antibiotic therapy

- ◆ Begin intravenous antibiotics as early as possible, and always within the first hour of recognizing severe sepsis (1D) and septic shock. (1B)
- ◆ Broad-spectrum: one or more agents active against likely bacterial/fungal pathogens and with good penetration into presumed source. (1B)
- ◆ Reassess antimicrobial regimen daily to optimize efficacy, prevent resistance, avoid toxicity, & minimize costs. (1C)
- ◇ Consider combination therapy in Pseudomonas infections. (2D)
- ◇ Consider combination empiric therapy in neutropenic patients. (2D)
- ◇ Combination therapy no more than 3-5 days and de-escalation following susceptibilities. (2D)
- ◆ Duration of therapy typically limited to 7–10 days; longer if response slow, undrainable foci of infection, or immunologic deficiencies. (1D)
- ◆ Stop antimicrobial therapy if cause is found to be non-infectious. (1D)

Source identification and control

- ◆ A specific anatomic site of infection should be established as rapidly as possible (1C) and within the first 6 hours of presentation. (1D)
- ◆ Formally evaluate patient for a focus of infection amenable to source control measures (eg: abscess drainage, tissue debridement). (1C)
- ◆ Implement source control measures as soon as possible following successful initial resuscitation. (1C)
- ◇ Exception: infected pancreatic necrosis, where surgical intervention best delayed. (2B)
- ◆ Choose source control measure with maximum efficacy and minimal physiologic upset. (1D)
- ◆ Remove intravascular access devices if potentially infected. (1C)

Fluid therapy

- ◆ Fluid-resuscitate using crystalloids or colloids. (1B)
- ◆ Target a CVP of ≥ 8 mmHg (≥ 12 mmHg if mechanically ventilated). (1C)
- ◆ Use a fluid challenge technique while associated with a hemodynamic improvement. (1D)
- ◆ Give fluid challenges of 1000 mL of crystalloids or 300–500 mL of colloids over 30 minutes. More rapid and larger volumes may be required in sepsis-induced tissue hypoperfusion. (1D)
- ◆ Rate of fluid administration should be reduced if cardiac filling pressures increase without concurrent hemodynamic improvement. (1D)

Vasopressors

- ◆ Maintain MAP ≥ 65 mmHg. (1C)
- ◆ Norepinephrine or dopamine centrally administered are the initial vasopressors of choice. (1C)
- ◆ Epinephrine, phenylephrine, or vasopressin should not be administered as the initial vasopressor in septic shock. (2C)
 - Vasopressin 0.03 units/min may be subsequently added to norepinephrine with anticipation of an effect equivalent to norepinephrine alone.
- ◇ Use epinephrine as the first alternative agent in septic shock when blood pressure is poorly responsive to norepinephrine or dopamine. (2B)
- ◆ Do not use low-dose dopamine for renal protection. (1A)
- ◆ In patients requiring vasopressors, insert an arterial catheter as soon as practical. (1D)

Inotropic therapy

- ◆ Use dobutamine in patients with myocardial dysfunction as supported by elevated cardiac filling pressures and low cardiac output. (1C)
- ◆ Do not increase cardiac index to predetermined supranormal levels. (1B)

Steroids

- ◇ Consider intravenous hydrocortisone for adult septic shock when hypotension responds poorly to adequate fluid resuscitation and vasopressors. (2C)
- ◇ ACTH stimulation test is not recommended to identify the subset of adults with septic shock who should receive hydrocortisone. (2B)
- ◇ Hydrocortisone is preferred to dexamethasone. (2B)
- ◇ Fludrocortisone (50 µg orally once a day) may be included if an alternative to hydrocortisone is being used which lacks significant mineralocorticoid activity. Fludrocortisone is optional if hydrocortisone is used. (2C)
- ◇ Steroid therapy may be weaned once vasopressors are no longer required. (2D)
- ◆ Hydrocortisone dose should be ≤300mg/day. (1A)
- ◆ Do not use corticosteroids to treat sepsis in the absence of shock unless the patient's endocrine or corticosteroid history warrants it. (1D)

Recombinant human activated protein C (rhAPC)

- ◇ Consider rhAPC in adult patients with sepsis-induced organ dysfunction with clinical assessment of high risk of death (typically APACHE II ≥ 25 or multiple organ failure) if there are no contraindications. (2B; 2C for post-operative patients)
- ◆ Adult patients with severe sepsis and low risk of death (eg: APACHE II ≤20 or one organ failure) should not receive rhAPC. (1A)

Blood product administration

- ◆ Give red blood cells when hemoglobin decreases to ≤7.0 g/dL (≤70 g/L) to target a hemoglobin of 7.0 – 9.0 g/dL in adults. (1B)
A higher hemoglobin level may be required in special circumstances (eg: myocardial ischemia, severe hypoxemia, acute hemorrhage, cyanotic heart disease, or lactic acidosis)
- ◆ Do not use erythropoietin to treat sepsis-related anemia. Erythropoietin may be used for other accepted reasons. (1B)
- ◇ Do not use fresh frozen plasma to correct laboratory clotting abnormalities unless there is bleeding or planned invasive procedures. (2D)
- ◆ Do not use antithrombin therapy. (1B)
- ◇ Administer platelets when: (2D)
 - counts are ≤5000/mm³ (5 X 10⁹/L) regardless of bleeding.
 - counts are 5000 to 30,000/mm³ (5–30 X 10⁹/L) and there is significant bleeding risk.
 - Higher platelet counts ≥ 50,000/mm³ (50 X 10⁹/L) are typically required for surgery or invasive procedures.

Mechanical ventilation of sepsis-induced acute lung injury (ALI)/ARDS

- ◆ Target a tidal volume of 6mL/kg (predicted) body weight in patients with ALI/ARDS. (1B)
- ◆ Target an initial upper limit plateau pressure ≤30cmH₂O. Consider chest wall compliance when assessing plateau pressure. (1C)
- ◆ Allow PaCO₂ to increase above normal, if needed, to minimize plateau pressures and tidal volumes. (1C)
- ◆ Positive end expiratory pressure (PEEP) should be set to avoid extensive lung collapse at end expiration. (1C)
- ◇ Consider using the prone position for ARDS patients requiring potentially injurious levels of FiO₂ or plateau pressure, provided they are not put at risk from positional changes. (2C)
- ◆ Maintain mechanically ventilated patients in a semi-recumbent position unless contraindicated. (1B)
 - ◇ Suggested target elevation 30 - 45 degrees. (2C)
- ◇ Noninvasive ventilation may be considered in the minority of ALI/ARDS patients with mild-moderate hypoxemic respiratory failure. The patients need to be hemodynamically stable, comfortable, easily arousable, able to protect/clear their airway, and expected to recover rapidly. (2B)
- ◆ Use a weaning protocol and a spontaneous breathing trial (SBT) regularly to evaluate the potential for discontinuing mechanical ventilation. (1A)
 - SBT options include a low level of pressure support with continuous positive airway pressure 5 cm H₂O or a T-piece.
 - Before the SBT, patients should:
 - be arousable
 - be hemodynamically stable without vasopressors
 - have no new potentially serious conditions
 - have low ventilatory and end-expiratory pressure requirement
 - require FiO₂ levels that can be safely delivered with a face mask or nasal cannula
- ◆ Do not use a pulmonary artery catheter for the routine monitoring of patients with ALI/ARDS. (1A)
- ◆ Use a conservative fluid strategy for patients with established ALI who do not have evidence of tissue hypoperfusion. (1C)

Sedation, analgesia, and neuromuscular blockade in sepsis

- ◆ Use sedation protocols with a sedation goal for critically ill mechanically ventilated patients. (1B)
- ◆ Use either intermittent bolus sedation or continuous infusion sedation to predetermined end points (sedation scales), with daily interruption/lightening to produce awakening. Re-titrate if necessary. (1B)
- ◆ Avoid neuromuscular blockers where possible. Monitor depth of block with train-of-four when using continuous infusions. (1B)

Glucose control

- ◆ Use IV insulin to control hyperglycemia in patients with severe sepsis following stabilization in the ICU. (1B)
- ◇ Aim to keep blood glucose ≤8.3 mmol/L (150 mg/dL) using a validated protocol for insulin dose adjustment. (2C)
- ◆ Provide a glucose calorie source and monitor blood glucose values every 1-2 hours (4 hours when stable) in patients receiving intravenous insulin. (1C)
- ◆ Interpret with caution low glucose levels obtained with point of care testing, as these techniques may overestimate arterial blood or plasma glucose values. (1B)

Renal replacement

- ◇ Intermittent hemodialysis and continuous veno-venous hemofiltration (CVVH) are considered equivalent. (2B)
- ◇ CVVH offers easier management in hemodynamically unstable patients. (2D)

Bicarbonate therapy

- ◆ Do not use bicarbonate therapy for the purpose of improving hemodynamics or reducing vasopressor requirements when treating hypoperfusion-induced lactic acidemia with pH ≥ 7.15. (1B)

Deep vein thrombosis (DVT) prophylaxis

- ◆ Use either low-dose unfractionated heparin (UFH) or low-molecular weight heparin (LMWH), unless contraindicated. (1A)
- ◆ Use a mechanical prophylactic device, such as compression stockings or an intermittent compression device, when heparin is contraindicated. (1A)
- ◇ Use a combination of pharmacologic and mechanical therapy for patients who are at very high risk for DVT. (2C)
- ◇ In patients at very high risk LMWH should be used rather than UFH. (2C)

Stress ulcer prophylaxis

- ◆ Provide stress ulcer prophylaxis using H₂ blocker (1A) or proton pump inhibitor (1B). Benefits of prevention of upper GI bleed must be weighed against the potential for development of ventilator-acquired pneumonia.

Consideration for limitation of support

- ◆ Discuss advance care planning with patients and families. Describe likely outcomes and set realistic expectations. (1D)