Management of Sepsis and Septic Shock

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Summary of the Clinical Problem
Sepsis results when the body’s response to infection causes life-threatening organ dysfunction. Septic shock is sepsis that results in tissue hypoperfusion, with vasopressor-requiring hypotension and elevated lactate levels. Sepsis is a leading cause of death, morbidity, and expense, contributing to one-third to half of deaths of hospitalized patients, depending on definitions. Management of sepsis is a complicated clinical challenge requiring early recognition and management of infection, hemodynamic issues, and other organ dysfunctions.

Characteristics of the Guideline Source
The guideline was developed by the SSC, with funding and governance from the SCCM and the ESICM (Table). Guideline committee members were from numerous specialties and included methods experts and a patient representative. A formal conflict of interest management policy was followed.

Evidence Base
The guideline committee used the GRADE method. Population, intervention, control, and outcomes questions were constructed; professional librarians assisted with evidence reviews. Although the 2016 revision of definitions for sepsis were published during the guideline development process, studies used for guideline evidence used earlier definitions of sepsis syndromes.

Benefits and Harms
The 2012 sepsis guidelines strongly recommended protocolized resuscitation with quantitative end points (early goal-directed therapy [EGDT]). Recommendations included specific goals for central venous pressure (CVP), mean arterial pressure (MAP), and central venous oxygen saturation and formed the basis of national quality and performance metrics. Since the 2012 guideline, substantial evolution has occurred in understanding the value of EGDT. Three key randomized trials enrolled patients presenting to the emergency department who had...
Sepsis with shock or hypoperfusion. In the PROCESS trial (n=1341 patients from 31 US institutions), protocol-based approaches did not reduce 60-day mortality vs usual care (19.5% vs 18.9%; relative risk [RR], 1.04; 95% CI, 0.82-1.31; P = .83).6 The similarly sized UK-based PROMISE7 and the ARISE trial8 from Australia and New Zealand both compared EGDT and usual care at 90 days and again found no difference in mortality (29.5% vs. 29.2%; RR, 1.01; 95% CI, 0.85-1.20; P = .90 and 18.6% vs 18.8%; RR, 0.98; 95% CI, 0.80-1.21; P = .90, respectively). Taken together, these trials suggest that while EGDT is safe, it is not superior to usual, nonprotocolized care. Usual care has also evolved since these trials to include more aggressive fluid resuscitation.9 In response, the 2016 guideline has removed standard EGDT resuscitation targets, instead recommending that sepsis-induced hypoperfusion be treated with at least 30 mL/kg of intravenous crystalloid given in 3 hours or less (Video 2). In the absence of the former static EGDT targets (eg, CVP), the guideline emphasizes frequent clinical reassessment and the use of dynamic measures of fluid responsiveness (eg, arterial pulse pressure variation), given evidence that dynamic measures predict fluid responsiveness better than static measures do.

Because infection causes sepsis, managing infection is perhaps the most critical component of sepsis therapy. Mortality increases even with very short delays of antimicrobials. To optimize the risk-benefit profile, the strategy of initial broad-spectrum therapy requires meticulous attention to antimicrobial stewardship, including early appropriate cultures and daily review to reduce or stop antimicrobials. Additionally, anatomic source control (eg, identifying infected central lines, pyelonephritis with ureteral obstruction, intestinal perforation) should occur as soon as is practical.

Discussion
The PROCESS,6 PROMISE,7 and ARISE8 trials have created substantial uncertainty in how to guide clinicians managing patients with sepsis and septic shock.9 When usual care is equivalent to EGDT, what is a clinician to do? The most significant update to the guideline reflects this shift in evidence: removing most specific EGDT endpoints and emphasizing frequent reevaluation and patient-specific tailoring of hemodynamic therapy. Even with a change in consensus definitions for sepsis,1 the guideline provides strong recommendations for a number of elements of standardized care, such as antimicrobial therapy, initial fluid volume, blood pressure goals, and vasopressor choice. Reflecting substantial consensus among experts, voting was by 75% of panel members with at least 80% agreement.

The guideline also provides a BPS for hospitals and health systems to develop formal sepsis performance improvement programs, given a suggestion of mortality benefit. Tools such as order sets, checklists, posters, reminder cards, and electronic medical record decision support may assist clinicians in early recognition and appropriate treatment of sepsis.10 Pediatric sepsis guidelines will be published separately, with a specific guideline for ventilation in ARDS expected in 2017.

Areas in Need of Future Study or Ongoing Research
The best approach for hemodynamic therapy for sepsis has become more uncertain as evidence has accumulated. This extends even to the degree to which clinicians should use intravenous fluids as a foundation for resuscitation in some patient groups. The guideline correctly identifies this as a key area for further research.

The best way to improve public health related to sepsis remains unsettled. For example, most US hospitals are required to report sepsis process measures. Collection of these data may be resource intense and may distract from other improvement efforts,5 inadvertently promote overtreatment or unnecessary testing, or delay nonsepsis diagnoses.3 At present, the international consensus definition of sepsis,1 the new guidelines,4 and CMS’s core measure requirements are unsynchronized. Thoughtful alignment would ensure meaningful reporting and improve patient outcomes.

Related Guidelines and Other Resources
UK National Institute for Health and Care Excellence (NICE)
Surgical Infection Society and Infectious Diseases Society of America (abdominal infections)
Infectious Diseases Society of America and American Thoracic Society (ventilator-associated pneumonia)
Sepsis guideline panelists (Video 4)