Sepsis: Early Recognition and Optimized Treatment

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Sepsis is a life-threatening condition caused by infection and represents a substantial global health burden. Recent epidemiological studies showed that sepsis mortality rates have decreased, but that the incidence has continued to increase. Although a mortality benefit from early-goal directed therapy (EGDT) in patients with severe sepsis or septic shock was reported in 2001, three subsequent multicenter randomized studies showed no benefits of EGDT versus usual care. Nonetheless, the early administration of antibiotics and intravenous fluids is considered crucial for the treatment of sepsis. In 2016, new sepsis definitions (Sepsis-3) were issued, in which organ failure was emphasized and use of the terms ‘systemic inflammatory response syndrome’ and ‘severe sepsis’ was discouraged. However, early detection of sepsis with timely, appropriate interventions increases the likelihood of survival for patients with sepsis. Also, performance improvement programs have been associated with a significant increase in compliance with the sepsis bundles and a reduction in mortality. To improve sepsis management and reduce its burden, in 2017, the World Health Assembly and World Health Organization adopted a resolution that urged governments and healthcare workers to implement appropriate measures to address sepsis. Sepsis should be considered a medical emergency, and increasing the level of awareness of sepsis is essential.

Keywords: Compliance; Mortality; Sepsis; Treatment

Introduction

Sepsis is a major cause of death from infection and represents a substantial healthcare burden, accounting for 6.2% of total hospital costs in the United States 2011. The estimated annual incidence of sepsis in the United States was 751,000 cases (3 cases/1,000 population) and the estimated number of deaths was 215,000. Recent large-scale epidemiological studies showed that the mortality rate of sepsis has decreased but its incidence continues to increase3,4. However, the true incidence of sepsis is likely to be underestimated.

On May 2017, the World Health Assembly (WHA) and World Health Organization (WHO) made sepsis a global health priority and adopted a resolution that urged the 194 United Nations Member States to improve the prevention, diagnosis, and management of sepsis. Accordingly, to improve patient outcomes, strategies that incorporate early recognition and timely management of sepsis in hospitals are being implemented5-8. In 2001, Rivers et al.9 reported the groundbreaking study on early-goal directed therapy (EGDT). However, three subsequent multicenter randomized controlled trials (RCTs) did not show that EGDT reduced the sepsis mortality rate compared to usual care10-12. Recently, new sepsis definitions were issued by the Society of Critical Care Medicine (SCCM) and the European Society of Intensive Care Medicine (ESICM) for screening and early identification. However, their benefits have yet to be validated by prospective studies13,14,15, and experts continue to place emphasis on the early administration of antibiotics and fluids for the initial resuscitation of patients.
with sepsis.

**Definitions and Early Identification of Sepsis**

1. Change in sepsis definitions

   The definition of sepsis has changed several times since 1992. The SCCM and ESICM revised the definition of sepsis and septic shock in 2016. The new definitions (Sepsis-3) focused on a dysregulated host response to infection and organ dysfunction. Sepsis is defined as infected patients with an increase of ≥2 Sequential Organ Failure Score (SOFA) points. Septic shock is defined as refractory hypotension requiring vasopressors with concurrent hyperlactemia (>2 mmol/L) despite adequate fluid resuscitation. Severe sepsis was excluded from the guidelines, and quick SOFA (qSOFA), instead of the systemic inflammatory response syndrome (SIRS), was adopted for screening purposes.

   The Sepsis-3 definitions were based on a large database and were the first to be tested in derivation and validation datasets. However, the definitions were not endorsed by some organizations and there are several issues associated with them. First, lactate was not retained in the sepsis definition. Hence, by the Sepsis-3 definitions, patients with an increased lactate level but no hypotension (or compensated septic shock) can be missed. In other words, we may miss patients in the early phase of sepsis. The prevalence of this phenotype (i.e., normotensive patients with hyperlactemia) was 26% in a previous multicenter trial. In the Sepsis-3 datasets, the prevalence of normotensive hyperlactemia (>4 mmol/L) was 9.9% but their mortality rate was not low (29.9%). Therefore, the validity of the Sepsis-3 definitions is suspect. Second, using the Sepsis-3 definitions, two components (the need for vasopressors and hyperlactemia) are needed concurrently to diagnose septic shock. That is, the lactate level is not a component of the definitions until the patient becomes hypotensive. Also, an infected patient with hypotension might not be considered to be in septic shock unless the lactate level was known. This implies that the utility of the Sepsis-3 definitions is limited in low-resource settings, where lactate levels are not frequently available. Therefore, further prospective studies are needed to demonstrate the validity of the Sepsis-3 definitions. Until then, it seems acceptable to use the pre-existing sepsis definitions.

2. Sepsis screening

   Sepsis screening is reportedly associated with a decreased mortality rate. The surviving sepsis campaign (SSC) guide-
lines of 2016, as well as those of 2012, emphasize routine screening of potentially infected patients who are likely to be septic to improve the early identification and treatment of sepsis. They recommend that hospitals should have a performance improvement program that involves early recognition and management of sepsis.

The SIRS criteria have, since 1992, been used to screen and identify sepsis patients. To diagnose sepsis, at least two of the four SIRS criteria must be met. However, because SIRS can be triggered by a variety of infectious and noninfectious causes, it is insufficiently sensitive, and certainly not specific for sepsis. Hence, some patients who satisfy the SIRS criteria may not have sepsis, and vice versa. In this context, the Sepsis-3 Task Force discarded the concept of SIRS and introduced, instead, qSOFA for sepsis screening. The qSOFA is a simplified version of the SOFA score that comprises only three variables, and patients with a qSOFA score of ≥2 should be considered for the possibility of sepsis (Figure 2). The qSOFA is a readily available bedside tool without laboratory tests, and has better specificity than qSOFA for identifying patients. Therefore, use of qSOFA may be premature, and the SIRS criteria are a sensitive and useful bedside tool for sepsis screening outside the ICU.

Clinical evidence indicates that patients with acute deterioration or sepsis manifest clinical signs or symptoms several hours before the condition worsens. Early warning scores, such as the Modified Early Warning Score (MEWS), Early Warning Scoring System (EWSS), or National Early Warning Score (NEWS), were developed to screen patients at high risk of deterioration. Although strong evidence based on robust data is lacking, these scores showed a trend toward improved outcomes and, when coupled with an outreach service (i.e., rapid response teams or medical emergency teams), it facilitates timely initiation of the optimal treatments upon recognition of septic patients. Although respiratory or cardiac problems were the most common trigger for activations of such outreach teams, one study showed that sepsis as a cause of activations accounted for 19.9%, and EGDT was undertaken in 22.7%. Interestingly, Churpek et al. compared several early warning scores, including qSOFA, among patients outside the ICU. qSOFA had a higher specificity and lower sensitivity than SIRS, MEWS, and NEWS for predicting in-hospital death or ICU transfer. The SIRS criteria (≥2) were more rapid than qSOFA for identifying patients. Therefore, use of qSOFA may be premature, and the SIRS criteria are a sensitive and useful bedside tool for sepsis screening outside the ICU.

**Early Treatments and Optimal Resuscitation**

The EGDT was designed for the early detection of sepsis and timely optimization of hemodynamic parameters by continuous monitoring of central venous oxygen saturation (ScvO₂ > 70%), central venous pressure (8–12 mm Hg), mean arterial pressure (MAP ≥ 65 mm Hg), and urine output (>0.5 mL/kg/h). This protocolized treatment, when administered to patients with severe sepsis or septic shock before admission to the ICU, reduced the incidence of multi-organ dysfunction and significantly decreased the in-hospital mortality rate compared with standard care.

However, three international multicenter trials (Protocolized Care for Early Septic Shock [ProCESS] and Australasian Resuscitation in Sepsis Evaluation [ARISE], and Protocolized Management in Sepsis [ProMISE]) did not show any significant survival benefit compared to usual care. In a meta-analysis of individual participants in the three RCTs, the EGDT did not result in better outcomes than usual care but was associated with increased hospitalization costs. Therefore, the EGDT concept was weakened in the 2016 guidelines. However, initial fluid resuscitation with crystalloids is still being emphasized, and patients in the usual care groups received a considerable volume of fluids in these three RCTs (Table 1). Application of balanced crystalloids significantly decreased the rates of all-cause mortality, persistent renal insufficiency, and new dialysis treatments, compared to saline. However, the 2016 guidelines emphasize, instead, the re-evaluation of volume status and tissue perfusion after the initial fluid resuscitation. This is because the persistence of a positive daily fluid balance over time was strongly associated with a higher mortality rate in patients with sepsis. In this regard, the guidelines have recommended either repeated assessments of vital signs, cardiopulmonary status, capillary refill time, pulse, and skin findings, or measurement of the two of followings: CVP, ScvO₂, bedside cardiovascular ultrasound, and dynamic assessment of fluid responsiveness with passive leg raise or fluid challenge.

A population-based study in the United States reported that early central vein catheterization was associated with a lower in-hospital mortality rate. However, the three RCTs of EGDT highlighted that there was no benefit of invasive hemodynamic monitoring involving CVP and ScvO₂ if initial fluids and adequate antibiotics were administered to septic patients in a timely manner. Accordingly, the 2016 SSC guidelines do not contain pre-specified treatment CVP and ScvO₂ targets. CVP does not reflect intravascular volume status precisely and is not predictive of the fluid response. Instead, repeated measurements of lactate (i.e., lactate clearance) enable evaluation of the responsiveness to initial resuscitation, and echocardiography is a noninvasive method of assessing the volume status in patients on mechanical ventilator support.
Table 1. Comparison of enrollment, treatment, and outcome of EGDT studies

|                     | EGDT | Control | EGDT | Protocol-based standard Therapy | EGDT | Usual care | EGDT | Usual care | EGDT | Usual care |
|---------------------|------|---------|------|---------------------------------|------|------------|------|------------|------|------------|
| Location            | United States | United States | Australia, New Zealand, Finland, Ireland, and Hong Kong | United Kingdom |
| Enrolled patients   | 130  | 133     | 439  | 446                              | 456  | 793        | 798  | 625        | 626  |            |
| Age, yr             | 67.1±17.4 | 64.4±17.1 | 60±16.4 | 61±16.1                         | 62±16.0 | 62.7±16.4 | 63.1±16.5 | 66.4±14.6 | 64.3±15.5 |
| APACHE II score (baseline) | 21.4±6.9 | 20.4±7.4 | 20.8±8.1 | 20.6±7.4                         | 20.7±7.5 | 15.4±6.5 | 15.8±6.5 | 18.7±7.1 | 18.0±7.1 |
| Arterial catheter insertion, % | Required | Required | Required | Required                       | Required | 91.4 | 76.3         | 74.2 | 622         |
| Central vein catheterization, % | Required | Required | 936  | 56.5                              | 57.9  | 90         | 61.9  | 92.1       | 509  |            |
| Initial lactate >4 mmol/L, % | 79   | 59       | 59.2  | 60.7                             | 46    | 46.5       | 65.4  | 637         |      |            |
| Total fluid within 6 hr, mL | 4,981±2,984 | 3,499±2,438 | 5,059 | 5,511                            | 4,362 | 4,479      | 4,304 | 4,100       | 4,074 |            |
| Fluid prior to randomization (at ED)* | -    | -       | 2,254±1,472 | 2,226±1,363 | 2,083±1,405 | 2,515±1,244 | 2,591±1,331 | 1,600 | 1,790       | (1,000–2,500) | (1,000–2,500) |
| Use of vasopressors within 6 hr, % | 27.4 | 30.3     | 54.9  | 52.2                             | 44.1  | 66.6       | 57.8  | 53.3        | 466  |            |
| RBC transfusion within 6 hr, % | 64.1 | 18.5     | 14.4  | 8.3                              | 7.5   | 13.6       | 7.0   | 8.8         | 3.8  |            |
| Length of ICU stay, day* | -    | -       | 5.1±6.3 | 5.1±7.1                         | 4.7±5.8 | 2.8 (1.4–5.1) | 2.8 (1.5–5.7) | 2.6 (1.0–5.8) | 2.2 (0.0–5.3) |
| Hospital mortality, % | 30.5 | 46.5     | 21.0  | 18.2                             | 18.9  | 14.5       | 15.7  | 23.6        | 24.6 |            |

*Mean±standard deviation or median (interquartile range).
EGDT: early goal-directed therapy; ProCESS: Protocolized Care for Early Septic Shock; ARISE: Australasian Resuscitation in Sepsis Evaluation; ProMISE: Protocolized Management in Sepsis; APACHE II: Acute Physiology and Chronic Health Evaluation II; ED: emergency department; RBC: red blood cell; ICU: intensive care unit.
Delayed administration of empirical antibiotics after sepsis identification increases the in-hospital mortality rate. Liu et al. recently reported that a 1-hour delay in antibiotic initiation was associated with an increased odds of in-hospital mortality among patients who received antibiotics within 6 hours. Therefore, the SSC guidelines recommend the intravenous administration of empiric antibiotics after obtaining blood culture results within 1 hour. Treatment with one or two broad-spectrum antibiotics and early de-escalation after clinical improvement or pathogen non-detection are recommended.

Early administration of vasopressors is associated with an increased survival rate in patients with septic shock and is a component of the 6-hour sepsis bundle. Thus, norepinephrine, as the first choice, should be administered early to maintain a MAP ≥65 mm Hg, when hypotension does not respond to initial fluid resuscitation. In an open-label RCT, targeting a MAP of 80–85 mm Hg rather than 65–70 mm Hg did not increase the survival rate of patients with septic shock. Thus, the target MAP should be determined according to the patients’ condition; a higher target may be needed for patients with chronic hypertension and a lower target for those with uncontrolled bleeding with trauma.

The SSC guidelines do not include a target heart rate for patients with septic shock. However, due to the many adverse effects of tachycardia, such as diastolic dysfunction and myocardial ischemia, the heart rate should be maintained within the normal range in patients with septic shock. Recently, Morelli et al. demonstrated that esmolol can be safely used to reduce the heart rate, which may facilitate accurate estimation of the incidence of sepsis, which can be used to improve performance and formulate future policies. A good example is the New York State sepsis registry. Beginning in 2014, all hospitals in New York State adopted sepsis protocols based on Rory’s regulations for the early diagnosis and treatment of sepsis. They are required to report their performance (compliance), as well as clinical information, to the New York State Department of Health. Surprisingly, after the onset of the initiative, the average rate of protocol compliance has increased progressively from 73.7% in the second quarter of 2014 to 84.7% of adult sepsis patients in the third quarter of 2016. This compliance is in contrast to Asian countries; a large multinational study of Asian ICUs reported rates of compliance with the resuscitation and management bundles of 7.6% and 3.5%, respectively.

The Core Outcome and Resource Evaluation (CORE) committee is a component of the Australia and New Zealand Intensive Care Society (ANZICS). All ICUs in Australia and New Zealand were invited to contribute to the ANZICS CORE registries in 1992. These registries have four registry domains: adult patients, pediatric patients, critical care resources, and central line-associated bloodstream infections. The CORE committee collects comprehensive data on various aspects of ICUs and patients to increase the level of awareness of the general public. However, despite the significant impact, public awareness is currently very low; a survey in 2009 reported that 88% of respondents had never heard the term “sepsis.” Since 2012, the Global Sepsis Alliance (GSA) has organized the annual World Sepsis Day, and many organizations or countries are undertaking sepsis awareness campaigns. In healthcare facilities, continuous training and education are also needed to increase the level of awareness of sepsis on the part of healthcare providers, who must understand that the condition is a real medical emergency.

Most estimates of sepsis are based on studies in high-income countries, and data on low- and middle-income (or resource-limited) countries are scarce. Therefore, increasing the awareness of sepsis is essential for controlling the sepsis burden in those countries. In South Korea, on “World Sepsis Day,” annual symposia and field events have taken place since 2012. However, further promotional or educational activities for the public, as well as support from the political leadership, are required to improve the situation.

2. Building a sepsis registry

Sepsis is frequently handled like a “garbage code” in the Global Burden of Disease Statistics. Most deaths due to sepsis are classified as their underlying infections, rather than sepsis itself. Therefore, the burden of sepsis is likely to be underestimated. Hence, improving the coding process for sepsis may facilitate estimation of the true burden.

Nationwide (or statewide) registry data collected prospectively can facilitate accurate estimation of the incidence of sepsis, which can be used to improve performance and formulate future policies. A good example is the New York State sepsis registry. Beginning in 2014, all hospitals in New York State adopted sepsis protocols based on Rory’s regulations for the early diagnosis and treatment of sepsis. They are required to report their performance (compliance), as well as clinical information, to the New York State Department of Health. Surprisingly, after the onset of the initiative, the average rate of protocol compliance has increased progressively from 73.7% in the second quarter of 2014 to 84.7% of adult sepsis patients in the third quarter of 2016. This compliance is in contrast to Asian countries; a large multinational study of Asian ICUs reported rates of compliance with the resuscitation and management bundles of 7.6% and 3.5%, respectively.

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Early recognition and treatment of sepsis

The five key elements of hour-1 bundle

1. Measure lactate level. Remeasure if initial lactate is >2 mmol/L.
2. Obtain blood cultures prior to administration of antibiotics.
3. Administer broad-spectrum antibiotics.
4. Begin rapid administration of 30 mL/kg crystalloid for hypotension or lactate >4 mmol/L.
5. Apply vasopressors if patient is hypotensive during or after fluid resuscitation to maintain MAP ≥65 mm Hg.

“Time zero” or “time of presentation” is defined as the time of triage in the Emergency Department, or if presenting from another care venue, from the earliest chart annotation consistent with all elements of sepsis (formerly severe sepsis) or septic shock ascertained through chart review.

Adapted from Levy et al. Crit Care Med. 2018;46:997-1000, with permission of Society of Critical Care Medicine. MAP: mean arterial pressure.

Table 2. Hour-1 surviving sepsis campaign bundle of care

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Conclusion

Prevention and early recognition of sepsis are of paramount importance until novel emerging drugs (or interventions) are demonstrated to be effective. Early application of the optimal treatments and improved compliance with sepsis bundles are pre-requisites for improving patients’ outcomes. The validity of the Sepsis-3 definitions and that of qSOFA need to be demonstrated in large-scale prospective trials.

Authors’ Contributions

Conceptualization: Kim HI, Park S. Methodology: Kim HI, Park S. Formal analysis: Kim HI, Park S. Original draft preparation: Kim HI, Park S. Review and editing: Kim HI, Park S. Approval of final manuscript: all authors.

Conflicts of Interest

No potential conflict of interest relevant to this article was reported.

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