



Sepsiste Bundle Uygulamaları

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Tıp Fakültesi

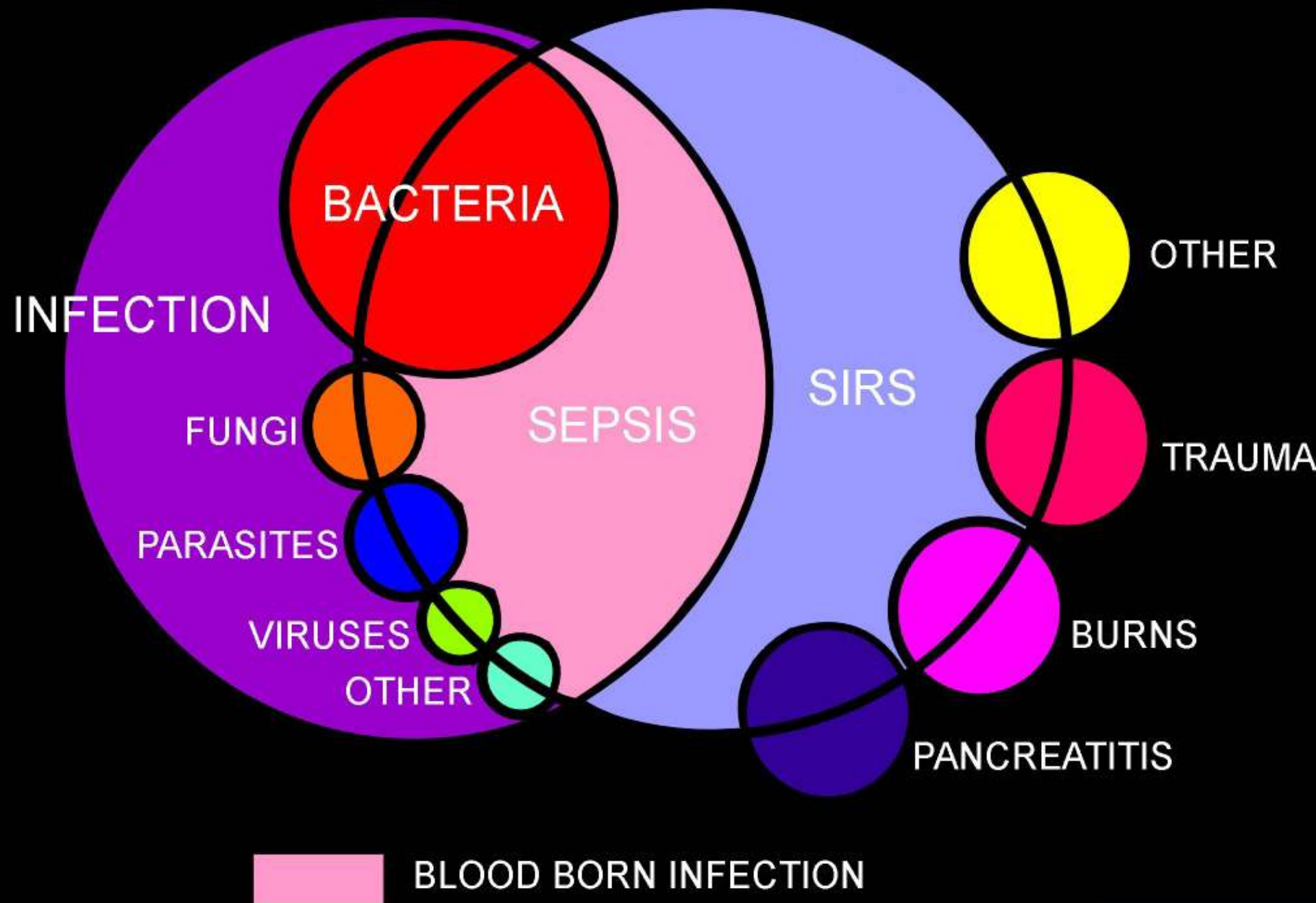
Enfeksiyon Hastalıkları ve Klinik Mikrobiyoloji Kliniği

Tanımlamalar

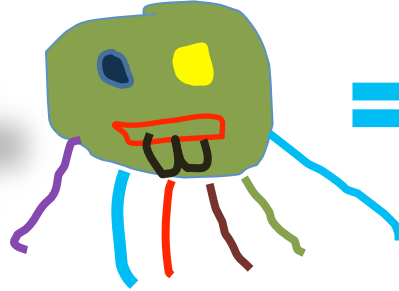
Bakteremi

SIRS

Sepsis



SIRS



Sepsis



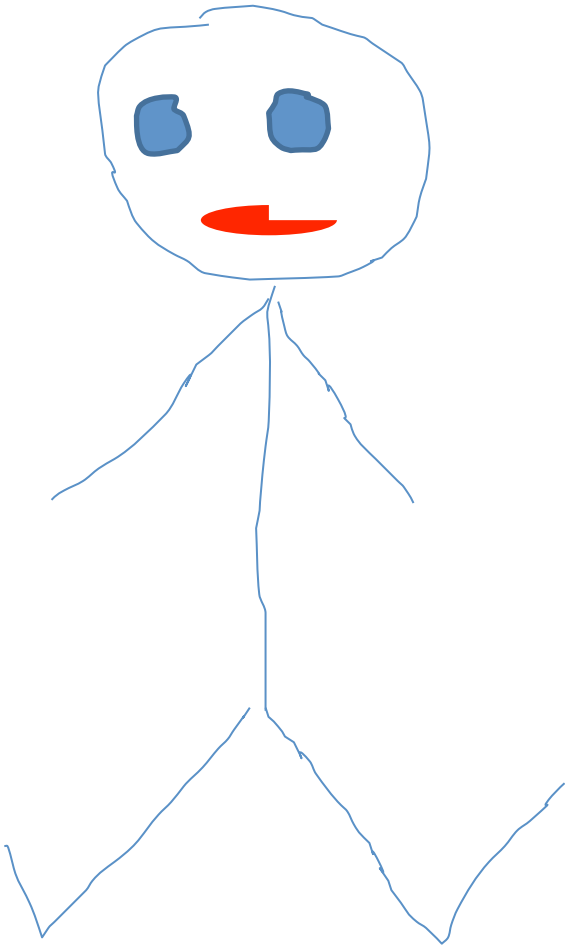
**Ađır Sepsis/
Sepsis
Sendromu**

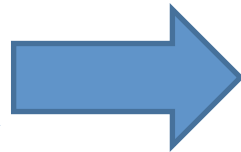
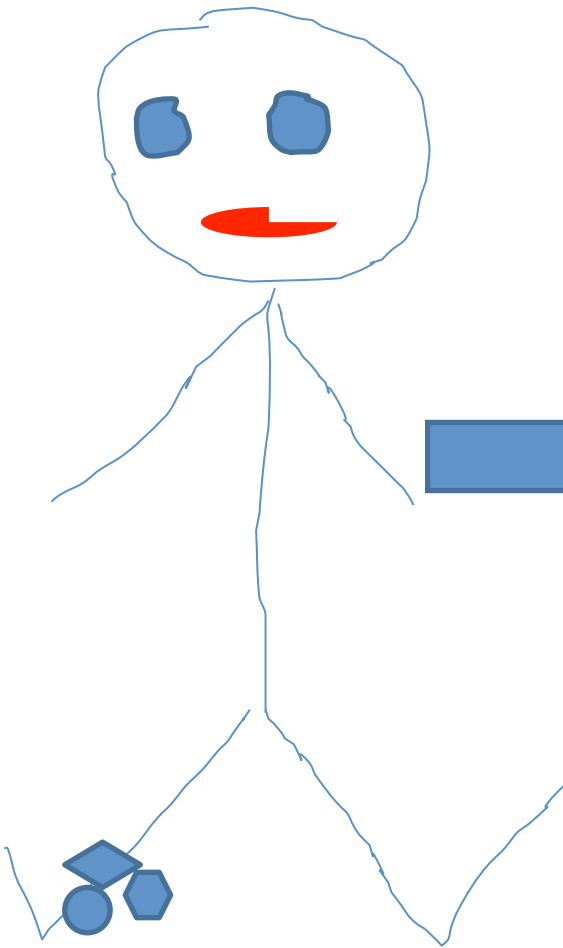


**Organ
yetmezliđi**

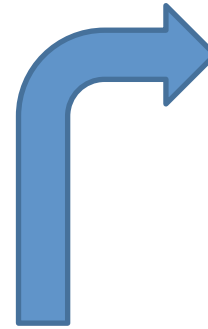
Hipoperfüzyon
Laktik asidoz
Oliguri
Mental durum
deđişikliđi

NORMAL HOST

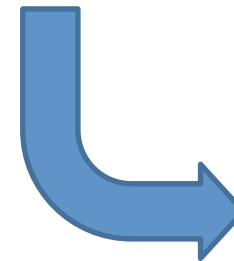




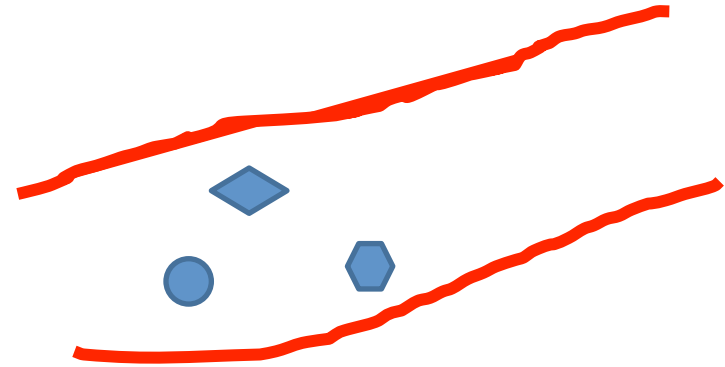
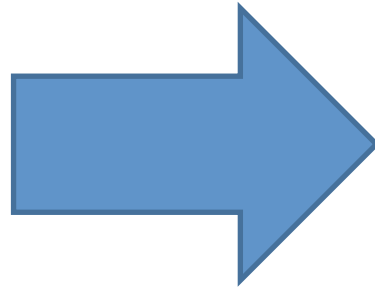
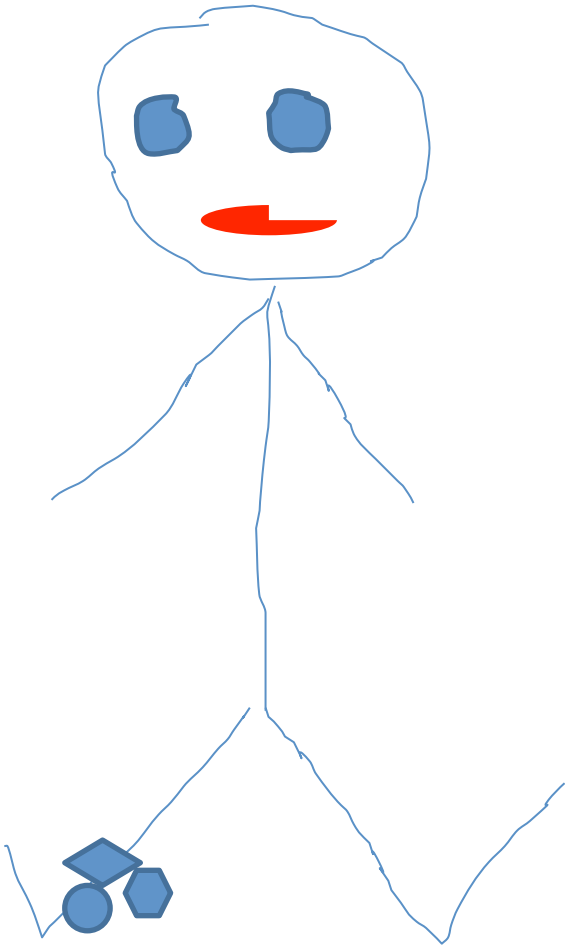
**interaction
between the
pathogen and
the host's
immune
system**



Host kills
bacteria



Bacteria kills
the host



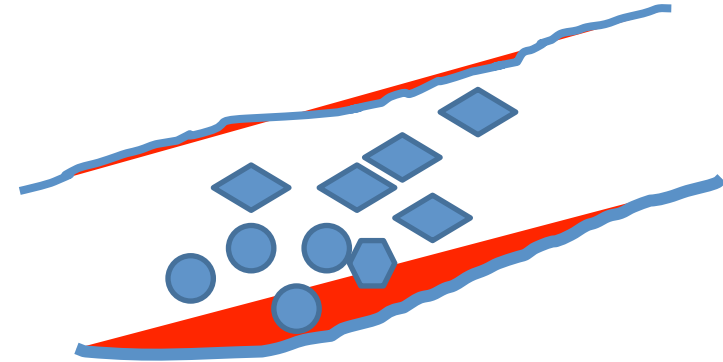
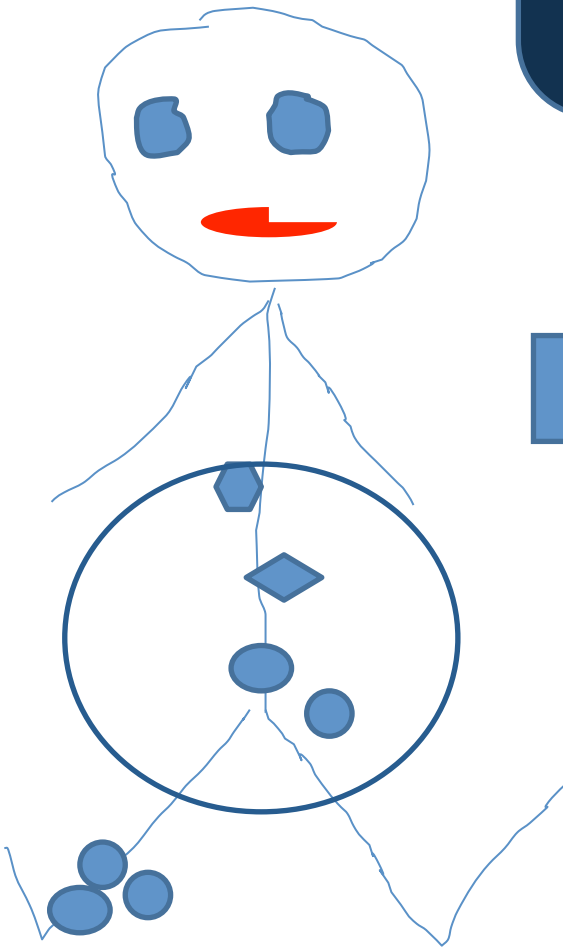
**Blood
vessel**

ENDOTOXINS

Cytokine release

LPS!!-----growth phases of the Gr neg bacteria
Cell lysis by host defence mechanism-antibiotics

**Peptidoglycan layer and
Teichoic acids** (Gram pos)



Blood vessel

EXOTOXINS

TSST!!- *S. aureus* and Gr A
Streptococcus

Large T cell and cytokine response

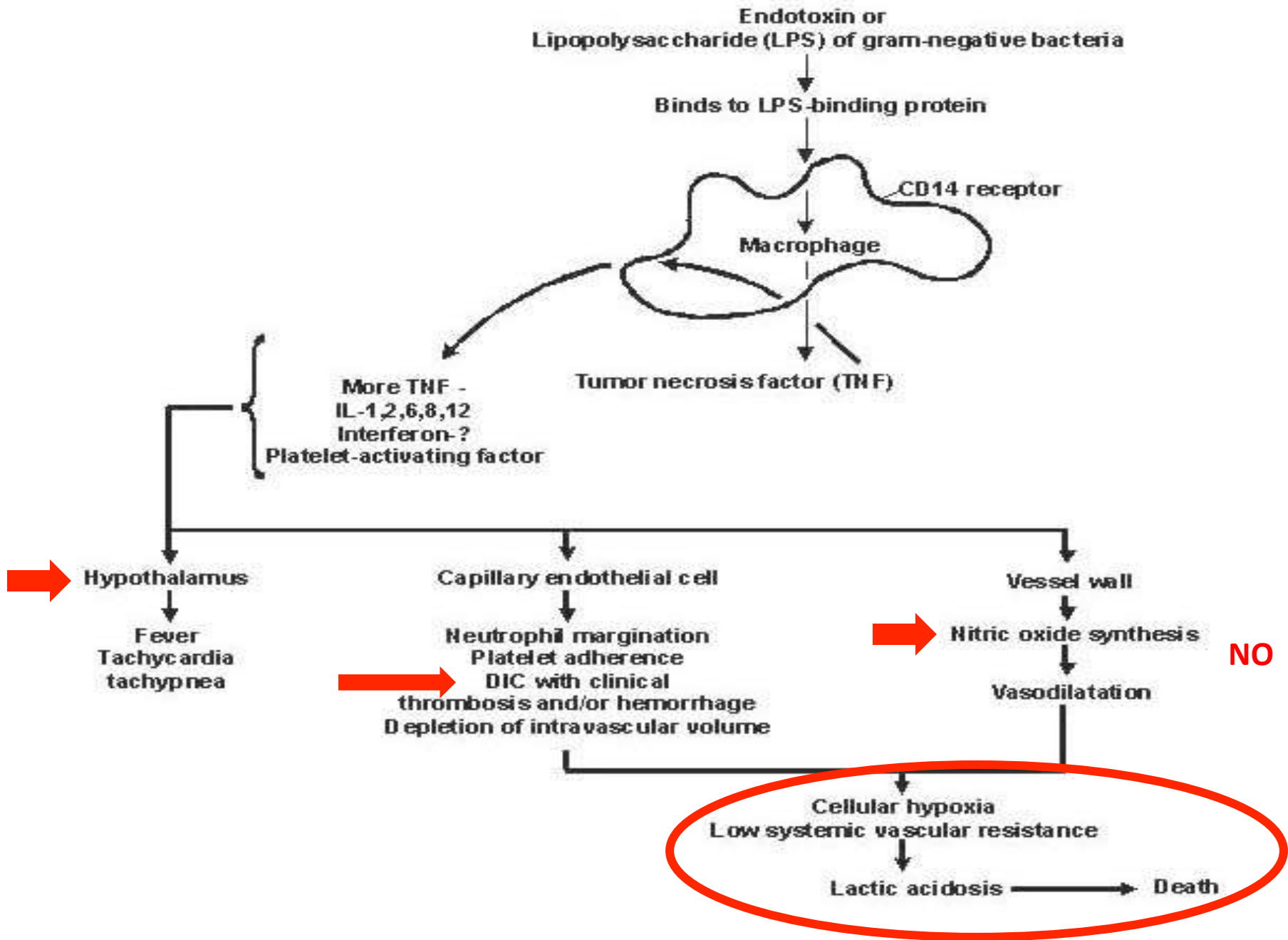
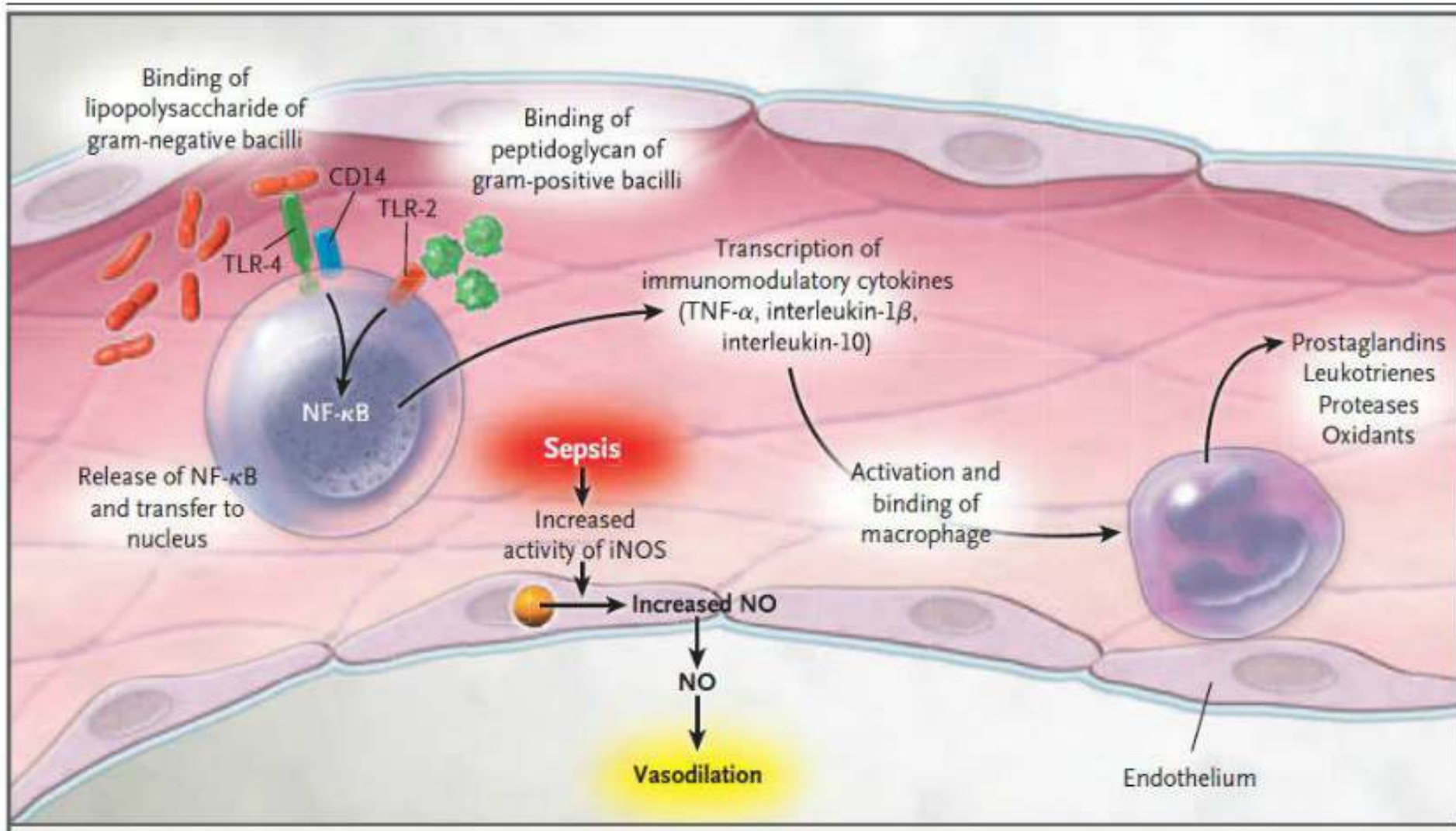


Diagram depicting the pathogenesis of sepsis and multiorgan failure. DIC = disseminated intravascular coagulation; IL = interleukin. Khalil A. Medscape 2014



Hypotension, the cardinal manifestation of sepsis, occurs via induction of nitric oxide (NO).

NO plays a major role in the hemodynamic alterations of septic shock, which is a hyperdynamic form of shock.

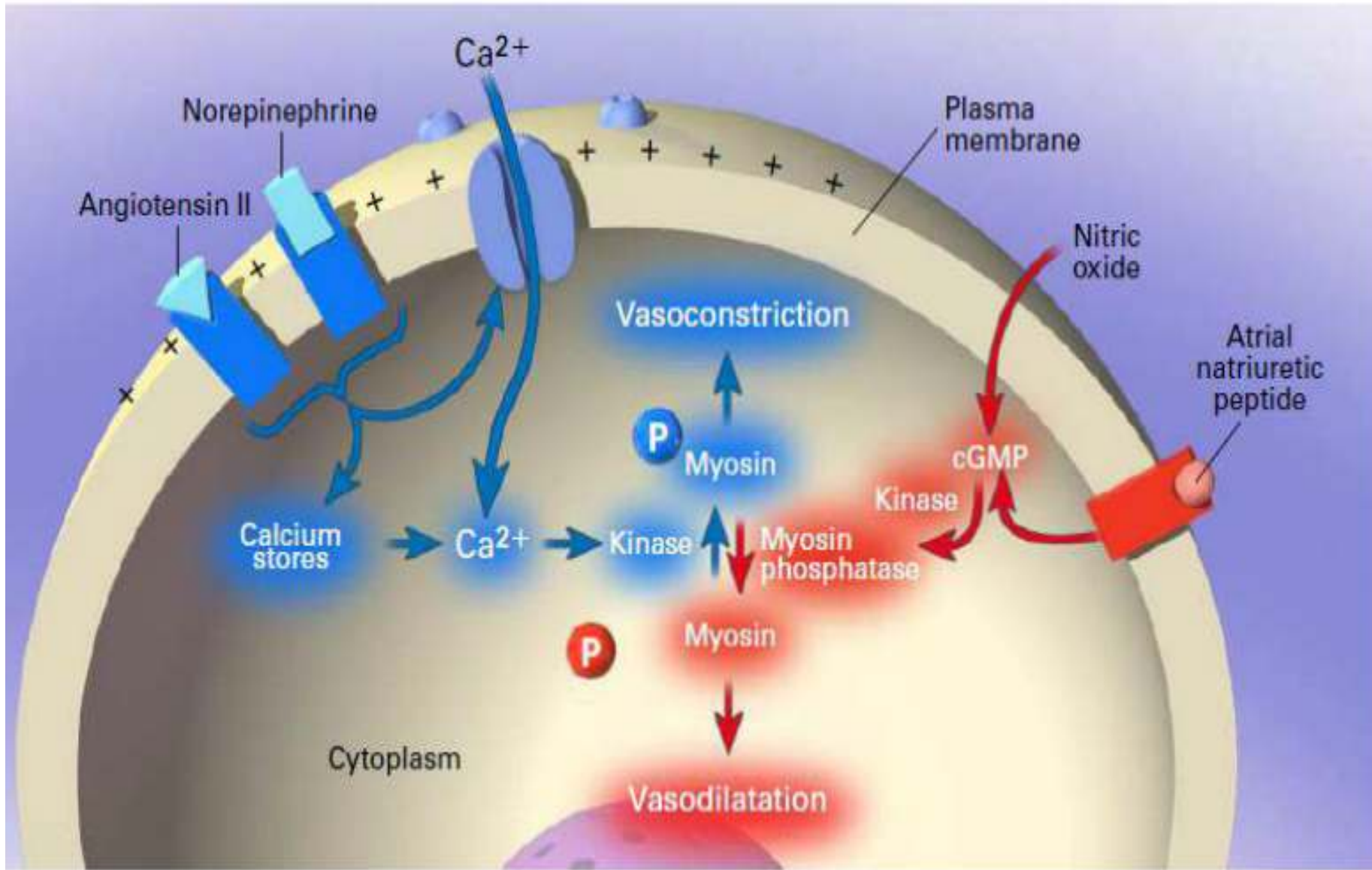


TABLE 1. Diagnostic Criteria for Sepsis**Infection, documented or suspected, and some of the following:**

General variables

- Fever ($> 38.3^{\circ}\text{C}$)
- Hypothermia (core temperature $< 36^{\circ}\text{C}$)
- Heart rate $> 90/\text{min}^{-1}$ or more than two sd above the normal value for age
- Tachypnea
- Altered mental status
- Significant edema or positive fluid balance ($> 20\text{ mL/kg}$ over 24 hr)
- Hyperglycemia (plasma glucose $> 140\text{ mg/dL}$ or 7.7 mmol/L) in the absence of diabetes

Inflammatory variables

- Leukocytosis (WBC count $> 12,000\ \mu\text{L}^{-1}$)
- Leukopenia (WBC count $< 4000\ \mu\text{L}^{-1}$)
- Normal WBC count with greater than 10% immature forms
- Plasma C-reactive protein more than two sd above the normal value
- Plasma procalcitonin more than two sd above the normal value

Hemodynamic variables

- Arterial hypotension (SBP $< 90\text{ mm Hg}$, MAP $< 70\text{ mm Hg}$, or an SBP decrease $> 40\text{ mm Hg}$ in adults or less than two sd below normal for age)

Organ dysfunction variables

- Arterial hypoxemia ($\text{Pao}_2/\text{Fio}_2 < 300$)
- Acute oliguria (urine output $< 0.5\text{ mL/kg/hr}$ for at least 2 hrs despite adequate fluid resuscitation)
- Creatinine increase $> 0.5\text{ mg/dL}$ or $44.2\ \mu\text{mol/L}$
- Coagulation abnormalities (INR > 1.5 or aPTT $> 60\text{ s}$)
- Ileus (absent bowel sounds)
- Thrombocytopenia (platelet count $< 100,000\ \mu\text{L}^{-1}$)
- Hyperbilirubinemia (plasma total bilirubin $> 4\text{ mg/dL}$ or $70\ \mu\text{mol/L}$)

Tissue perfusion variables

- Hyperlactatemia ($> 1\text{ mmol/L}$)
- Decreased capillary refill or mottling

WBC = white blood cell; SBP = systolic blood pressure; MAP = mean arterial pressure; INR = international normalized ratio; aPTT = activated partial thromboplastin time.

Diagnostic criteria for sepsis in the pediatric population are signs and symptoms of inflammation plus infection with hyper- or hypothermia (rectal temperature $> 38.5^{\circ}$ or $< 35^{\circ}\text{C}$), tachycardia (may be absent in hypothermic patients), and at least one of the following indications of altered organ function: altered mental status, hypoxemia, increased serum lactate level, or bounding pulses.

Adapted from Levy MM, Fink MP, Marshall JC, et al: 2001 SCCM/ESICM/ACCP/ATS/SIS International Sepsis Definitions Conference. *Crit Care Med* 2003; 31: 1250-1256

Surviving Sepsis Campaign

International
Guidelines for
Management of
Severe Sepsis and
Septic Shock: 2012

Critical Care Medicine
2013 Feb;41(2):580-637



Surviving Sepsis Campaign > Guidelines

Guidelines

The third edition of "Surviving Sepsis Campaign: International Guidelines for Management of Severe Sepsis and Septic Shock: 2012" appeared in the February 2013 issues of *Critical Care Medicine* and *Intensive Care Medicine*.

 [Surviving Sepsis Campaign: International Guidelines for Management of Severe Sepsis](#)

 [Complete Implementation and Improvement Guide](#)

Tables summarizing the recommendations can be a useful tool in clinical settings.

 [Initial Resuscitation and Infection Issues](#)

 [Hemodynamic Support and Adjunctive Therapy](#)

 [Other Supportive Therapy of Severe Sepsis](#)

 [Special Considerations in Pediatrics](#)

SSC Statements

Guideline Translations

The SSC Executive Committee has released an update based on the release of the ProCESS and ARISE trials:

HEDEF
5 yılda sepsise
bağlı
mortaliteyi
%25 azaltmak

History

Campaign's history Since its inception in 2002 the Campaign has achieved several key milestones.

Original Stated Goal of Campaign

To reduce mortality from sepsis by 25% in 5 years (that translates to 2009 from the date of publication of the first set of guidelines) via a 7-point agenda including:

- Building awareness of sepsis
- Improving diagnosis
- Increasing the use of appropriate treatment
- Educating healthcare professionals
- Improving post-ICU care
- Developing guidelines of care
- Implementing a performance improvement

Phase I

Phase II

Phase III

Phase IV

Phase I: Development of Awareness of Scope of

Late 2001/Early 2002

- Separate from Campaign activities, a sepsis conference was held in 2001. Updates to criteria established in 1991. Le Gall et al. Conference. *Crit Care Med.* 2003 31(4):125
- Campaign was formed by the Society of Critical Care Medicine and launched at the ESICM Annual Meeting in 2002. On to the "Barcelona Declaration."
- Steering committee formed with 3 representatives

January 2003

- Executive committee meeting held in Amsterdam

February 2003

7 Nokta

1. Farkındalığı arttırma
2. Tanıyı arttırma
3. Uygun tedavi
4. Sağlık personelinin eğitimi
5. Post-YBU bakımı geliştirme
6. Kılavuzların geliştirilmesi
7. Performans geliştirici program

- Bundles ▶
- Data Collection ▶
- Resources ▶
- Implement/Improve ▶
- Contact ▶

- Building awareness of sepsis
- Improving diagnosis
- Increasing the use of appropriate treatment
- Educating healthcare professionals
- Improving post-ICU care
- Developing guidelines of care
- Implementing a performance improvement program

“Implementing the Surviving Sepsis Campaign”
2005

- Phase I
- Phase II
- Phase III
- Phase IV

Phase III: Guideline Implementation, Behavior Change, and Data Collection

September 2003

- The Surviving Sepsis Campaign initiated a partnership to improve techniques to treatment of sepsis. The
- Education initiatives continue at critical care conferences

January 2004

...ned to determine direction
...ergency Physicians imp

“Surviving Sepsis Campaign Guidelines”, 2nd ed, Critical Care Medicine and Intensive Care Medicine
2008

“Surviving Sepsis Campaign Guidelines”
published along with revised bundles
2012

...ean clinicians at Mediterranean Critical Care School and international representatives gathered to begin

...s developed and distributed by SCCM in North America and ESICM in Europe
... tool initiated

...edicine including background papers for all guidelines published.

...d to support collaboration in data collection and performance improvement throughout the

Management of Severe Sepsis

Initial
Resuscitation

Diagnosis

Antibiotic
Therapy

Source
Control

Fluid Therapy

Vasopressors

Corticosteroids

Blood Product
Administration

Glucose
Control

Bicarbonate
Therapy

Surviving Sepsis
Campaign

Sepsis Guidelines 2012

Surviving Sepsis Campaign Bundles

SURVIVING SEPSIS CAMPAIGN CARE BUNDLES

TO BE COMPLETED WITHIN 3 HOURS:

- 1) Measure lactate level
- 2) Obtain blood cultures prior to administration of antibiotics
- 3) Administer broad spectrum antibiotics
- 4) Administer 30 mL/kg crystalloid for hypotension or lactate ≥ 4 mmol/L

TO BE COMPLETED WITHIN 6 HOURS:

- 5) Apply vasopressors (for hypotension that does not respond to initial fluid resuscitation) to maintain a mean arterial pressure (MAP) ≥ 65 mm Hg
- 6) In the event of persistent arterial hypotension despite volume resuscitation (septic shock) or initial lactate ≥ 4 mmol/L (36 mg/dL):
 - Measure central venous pressure (CVP)*
 - Measure central venous oxygen saturation (ScvO₂)*
- 7) Remeasure lactate if initial lactate was elevated*

*Targets for quantitative resuscitation included in the guidelines are CVP of ≥ 8 mm Hg, ScvO₂ of $\geq 70\%$, and normalization of lactate.

Laktat!
Kültür!
Antibiyotik!
Sıvı!

- F- Fluids, (Give and measure).
- O- Give Oxygen (If not contraindicated)
- C- Culture
- A- Antibiotics
- L- Lactate

1) Sepsise baęlı doku hipoperfüzyonlu hastalarda destek miktarı (başlangıç sıvı desteęine rağmen devam eden hipotansiyon veya kan laktat konsantrasyonunun > 4 mmol/L olması durumunda)

İlk 6 saatlik resüsitasyonda hedefler:

- a) CVP 8–12 mmHg
- b) MAP \geq 65 mm Hg
- c) ScvO₂ 70% veya 65% (1C).

2) Laktat düzeyi yüksek hastalarda hedef resüsitasyon laktatı normalize etmek (2C)

Surviving Sepsis Campaign



Updated Bundles in Response to New Evidence

Bundles

The Surviving Sepsis Campaign Bundles are the core of the sepsis improvement efforts. Using "bundles" simplifies the complex processes of the care patients with severe sepsis. A bundle is a selected set of elements of care that are evidence-based practice guidelines that, when implemented as a group, have an effect on outcomes beyond implementing the individual elements alone.

Bundle Implementation

Each hospital's sepsis protocol may be customized, but it must meet the standards created by the bundle. Enhancing reliability of these bundle elements allows teams to focus on aspects of the changes they are implementing to create a reliable system that achieves the goal of 25 percent reduction in mortality due to sepsis called for by the Surviving Sepsis Campaign.

Surviving Sepsis Campaign Bundles - Revised 4/2015

TO BE COMPLETED WITHIN 3 HOURS

- 1) Measure lactate level
- 2) Obtain blood cultures prior to administration of antibiotics
- 3) Administer broad spectrum antibiotics
- 4) Administer 30 ml/kg crystalloid for hypotension or lactic acidemia

*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 generation consistent with all elements of severe sepsis or septic shock ascertained through chart review.

TO BE COMPLETED WITHIN 6 HOURS

- 2) Apply vasopressors for hypotension that does not respond to initial fluid resuscitation) to maintain a mean arterial pressure (MAP) ≥ 65 mm Hg
- 4) In the ward or intensive care unit (ICU) administer blood fluid administration (MAP ≥ 65 mm Hg) or fluid administration ≥ 4 L (total), re-evaluate volume status using reflexion and document findings according to Table 1.

Revised
4/2015

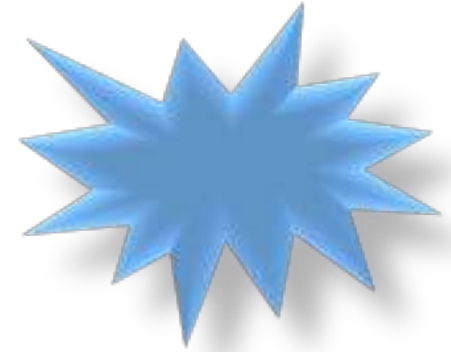
TO BE COMPLETED WITHIN 3 HOURS OF TIME OF PRESENTATION*:

1. Measure lactate level
2. Obtain blood cultures prior to administration of antibiotics
3. Administer broad spectrum antibiotics
4. Administer 30ml/kg crystalloid for hypotension or lactate ≥ 4 mmol/L

* *"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 severe sepsis or septic shock ascertained through chart review.*

3 saat içinde

1. Laktat seviyesini ölç
2. Antibiyotik öncesi KK al
3. Geniş spektrumlu antibiyotik başla
4. Hipotansiyon veya laktat ≥ 4 mmol/L ise 30ml/kg kristaloid başla



SIVI tedavisi

TO BE COMPLETED WITHIN 6 HOURS OF TIME OF PRESENTATION:

5. Apply vasopressors (for hypotension that does not respond to initial fluid resuscitation) to maintain a mean arterial pressure (MAP) ≥ 65 mmHg
6. In the event of persistent hypotension after initial fluid administration (MAP < 65 mm Hg) or if initial lactate was ≥ 4 mmol/L, re-assess volume status and tissue perfusion and document findings according to Table 1.
7. Re-measure lactate if initial lactate elevated.



6 saat içinde

5. **MAP ≥ 65 mmHg olacak şekilde vasopressor başla** (başlangıç sıvı tedavisine yanıt vermeyen sıvı resusitasyonunda)

6. **Başlangıç laktat düzeyi ≥ 4 mmol/L veya başlangıç sıvı tedavisine rağmen persistan hipotansiyon** varsa volüm durumunu ve doku perfüzyonunu tekrar gözden geçir, Tablo 1 e göre bulguları kaydet

7. Başlangıç laktat düzeyi yüksekse tekrar laktat düzeyini ölç

TABLE 1

DOCUMENT REASSESSMENT OF VOLUME STATUS AND TISSUE PERFUSION WITH:

EITHER

- Repeat focused exam (after initial fluid resuscitation) by licensed independent practitioner including vital signs, cardiopulmonary, capillary refill, pulse, and skin findings.

OR TWO OF THE FOLLOWING:

- Measure CVP
- Measure ScvO₂
- Bedside cardiovascular ultrasound
- Dynamic assessment of fluid responsiveness with passive leg raise or fluid challenge

Of note, the 6-hour bundle has been updated; the 3-hour SSC bundle is not affected.

While no suggestion of harm was indicated with use of a central line in any trial, and published evidence shows significant mortality reduction using the original SSC bundles (5), the committee has taken a prudent look at all current data and, despite weaknesses as in all studies, determined the above bundles to be the appropriate approach at this time.

**SBP <90 mmHg or MAP <65 mmHg
after 20-30cc/kg crystalloid IVF
-OR-
Lactate >4 mmol/L regardless of
blood pressure**

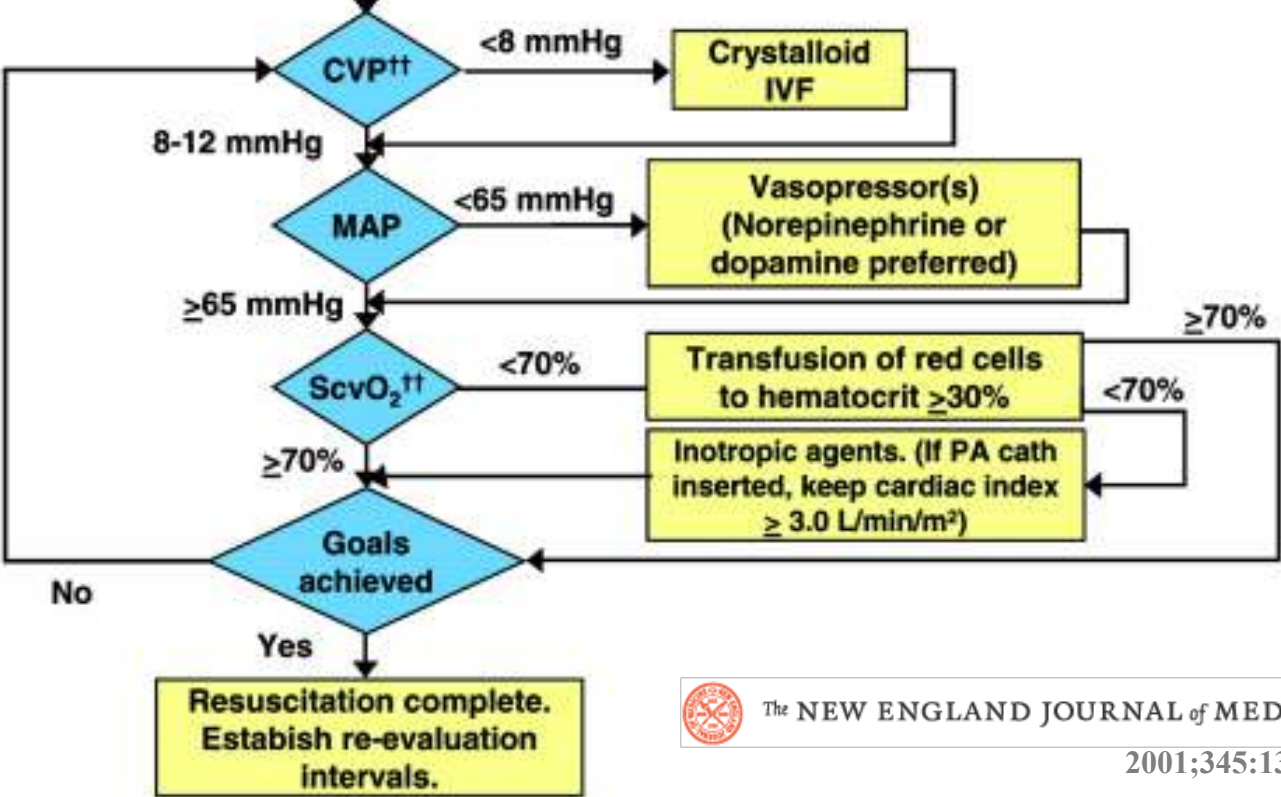
Supplemental oxygen ± endotracheal intubation and mechanical ventilation (if necessary)

Perform central venous catheterization while continuing crystalloid IVF resuscitation (250-1000 ml boluses)

Critical care consultation

This protocol should be initiated within ONE hour of presentation (or severe sepsis identification) and continued for at least the first SIX hours of resuscitation.

†† If PA catheter is inserted, PCWP target of 12-15 mmHg replaces CVP target, and SvO₂ ≥ 70% replaces ScvO₂ target.



SEPTIC SHOCK CLINICAL PATHWAY

Room # _____ ICU admission Date: _____ Time: _____

Please complete the following:

- ED Triage: Date: _____ Time: _____
- Septic Shock* diagnosis (Time Zero): Date: _____ Time: _____
- Patient transferred from (unit or hospital): _____
- Patient was identified as having severe sepsis or septic shock: ED Floor ICU Admission During ICU Stay
- Decision to move to comfort care in first 24 hours after diagnosis Yes No
- ICU discharge: Date: _____ Time: _____
- Discharge status: Alive Expired Attending physician at time of diagnosis: ED _____ ICU _____

*Septic Shock (Time Zero) defined as:
SBP less than 90mmHg or 40mmHg
decrease from baseline after 30ml/kg fluid
bolus, or requires vasopressors or initial
lactic acid is greater than or equal to 4mEq/L

Date _____ to _____ 0-1 Hours	Decision Grid	Date _____ to _____ 1-6 Hours	Date _____ to _____ 6-24 Hours	Date _____ to _____ 24-72 Hours
<input type="checkbox"/> Initial Labs: serum lactic acid, additional labs as ordered by physician _____ Serum lactic acid drawn Time _____ Yes No Blood Cultures X 2 Time 1: _____ Time 2: _____ <input type="checkbox"/> Other Cultures: <input type="checkbox"/> Establish IV access <input type="checkbox"/> Volume resuscitate: initial 30ml/kg over 1 hour or as fast as possible then additional boluses as needed per order _____ Time initial fluid bolus completed to resolve hypotension <input type="checkbox"/> Broad Spectrum Antibiotic-start after obtain blood culture (see Inonet under Pharmacy Guide to Antimicrobial Therapy) Yes No Was a new antibiotic initiated for this episode of septic shock? _____ Time antibiotic hung <input type="checkbox"/> Source Control	Yes No Is patient hypotensive after initial fluid bolus? Yes No Did patient require vasopressor(s)? If YES to either, continue to next column (Septic Shock Bundle) Yes No Is lactic acid greater or equal to 4 mEq/L? Yes No Is there evidence of additional organ dysfunction besides elevated lactic acid? If YES to lactic acid & additional organ dysfunction, please continue to next column (Septic Shock Bundle) If NO to additional organ dysfunction, please continue below: <input type="checkbox"/> Maintain MAP greater than or equal to 65 mmHg <input type="checkbox"/> Maintain U/O of 0.5 ml/kg/hour <input type="checkbox"/> Consider arterial line insertion <input type="checkbox"/> Monitor Stroke Volume & Stroke Volume Variation to guide fluid resuscitation <input type="checkbox"/> Ensure decrease in lactic acid x3 or normalization x2 within 12 hours • Begin next column (Septic Shock Bundle) if additional organ dysfunction occurs &/ or patient remains/becomes hypotensive after initial 30 ml/kg fluid bolus	Septic Shock Bundle Resuscitation Goals Yes No CVP placed If no, why? _____ _____ Time CVP placed (record first CVP reading prior to x-ray confirmation) Record the FIRST TIME the following is achieved: _____ Time CVP 8-12 mmHg on vent 12-15 mmHg _____ Time MAP greater than or equal to 65 mmHg _____ Time SCVO ₂ greater than 70%: mixed venous greater than or equal to 65% _____ Time Optimized stroke volume (optional) Yes No Assess for risk factors for abdominal compartment syndrome (fluid resuscitation greater than 5 L in 24 hours or less) _____ Repeat lactic acid every 4-6 hours	Septic Shock Bundle Yes No Is patient on vasopressor at greater than 6 hours Yes No Consider Vasopressin for refractory septic shock In patients with acute lung injury or ARDS; Yes No Patient on mechanical ventilator _____ PaO ₂ / FiO ₂ ratio Yes No Is tidal volume 6ml/kg of ideal body weight in first 24 hours? Yes No Are the static or plateau inspiratory pressures less than 30cmH ₂ O in first 24 hours?	<input type="checkbox"/> Confirm Infectious Source <input type="checkbox"/> Re-assess need for broad spectrum antibiotics based on culture reports. Yes No Was there an organism identified? Yes No If YES, was the organism sensitive to the initial antibiotic? <input type="checkbox"/> Discontinue Vancomycin if appropriate <input type="checkbox"/> Re-evaluate need for invasive lines and tubes <input type="checkbox"/> Nutrition Therapy <input type="checkbox"/> Progress Mobility
Nurse				
Nurse				
Physician				
Signature, Date & Time				

L

LEARN

E

ESTABLISH

A

ASK

D

DEVELOP

E

EDUCATE

R

REMEDiate

Examples Of Effective Teams

Example 1: Effective Work Team

Aim: Diagnose patients with severe sepsis or septic shock in the emergency department (ED) within 2 hours of triage

Core Working Team: The overall core team must be interdisciplinary and must include, at a minimum:

- ED physician
- Triage nurse
- Staff nurse
- Laboratory technician
- Laboratory supervisor
- Admissions clerk

Additional team members may include:

- Critical care medicine (CCM) physician
- House officer
- ICU charge nurse
- Infectious disease physician

Examples of Effective Aim Statements:

- Time from ED triage to presumptive diagnosis of severe sepsis is less than 2 hours
- Time from ED triage to all patients' meeting severe sepsis criteria having a serum lactate is less than 3 hours
- Time from ED triage to appropriate antibiotics given is less than 1 hour
- If hypotensive or if lactate > 4.0 mmol, immediate fluid resuscitation is started (at least 30 mL/kg normal saline or lactated ringers solution within 1 hour)
- If MAP < 65 mmHg and not responsive to adequate (at least 30 mL/kg) fluid resuscitation, vasopressors are started immediately
- If blood pressure or serum lactate not responsive to fluid, a central venous pressure monitor is instituted within the first 6 hours

Surviving Sepsis
Campaign

The logo for the Surviving Sepsis Campaign features the text "Surviving Sepsis" in a light green font and "Campaign" in a blue font below it. To the right of the text is a graphic consisting of several blue dots of varying sizes arranged in a curved pattern.



I M P R E S S

INTERNATIONAL **M**ULTICENTRE **P**REVALENCE **S**TUDY ON **S**EPSIS

Surviving Sepsis Campaign

An International **Multicentre **P**revalence **S**tudy on **S**epsis.**



IMPRESS

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THE PROTOCOL



REGISTER
TO PARTICIPATE



QUESTIONS &
ANSWERS



NATIONAL
COORDINATORS
LIST



STUDY
CRF



Hello: test test



Menu

Home

Users Configuration

Patient data

Patients

Historical

FAQ

Logout

This Project is endorsed by:

Surviving Sepsis
Campaign



The Intensive Care Society

Society of
Critical Care Medicine
The Intensive Care Professionals

CRF Document Eng.



Impress.pdf

User Document:



Impress_User
Manual.pdf



Powered By EDICS Platform



IMPRESS STUDY

International Multicentre PREvalence

Study on Sepsis

Turkey, USA, UK

Brezilya, Çek Cumhuriyeti, Danimarka

Yunanistan, Rusya

.....

The screenshot shows an email client interface. At the top, there are search buttons for 'Search Mail' and 'Search Web'. Below that, there are tabs for 'Deleted' and 'Calendar'. The main content area shows an email with the following details:

- From:** "Lori Harmon" <lharmon@sccm.org>
- To:** "Sepsisgroups@lists.sepsisgroups.org" <Sepsisgroups@lists.sepsisgroups.org>
- Subject:** [Sepsis Groups] IMPRESS Study update
- Date:** Monday, September 22, 2013 11:09 PM

The body of the email contains the following text:

All,

Enrolled countries for the [IMPRESS](#) Point Prevalence study has now reached 79 with 601 sites registered. Leading are the US with 92 sites, UK with 76, Spain and Italy at 50, Brazil at 21, Italy at 20 with Belgium, Czech Republic, Denmark, Turkey, Greece and Russia close in numbers. Dr. Rhodes reported this morning that the study is now registered on the NIH site <http://www.clinicaltrials.gov> use the search tag NCT01943747 if you are interested in locating it. Lets keep working together to get more sites enrolled. We are shooting for 1000 to participate in this important November 7th event.

Lori

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Society of Critical Care Medicine
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D +1-847-493-6403
F +1-847-493-6428
lharmon@sccm.org
<http://www.sccm.org>

R683

Publication Only

Clinical ID: Community-acquired infections including CAP, sepsis, STD,...

A survey of physicians' knowledge on evaluation of sepsis bundles: a multi-centre study

F. Civelek Eser¹, Z. Kocak Tufan¹, E. Vudali², A. Batirel³, B. Kayaaslan⁴, A. Tanrici Bastug⁴, D. Eray⁵, **V. Turhan**⁵, F. Duygu⁶, D. Tok⁷, S. Altun⁸

¹Infectious Disease and Clinical Microbiology, Yildirim Beyazit University Ankara Ataturk Training, Ankara, Turkey ; ²Infectious Disease and Clinical Microbiology, Ankara Training and Research Hospital, Ankara, Turkey ; ³Infectious Disease and Clinical Microbiology, Dr. Lutfi Kirdar Kartal Training and Research Hospital, Istanbul, Turkey ; ⁴Infectious Disease and Clinical Microbiology, Numune Training and Research Hospital, Ankara, Turkey ; ⁵Infectious Disease and Clinical Microbiology, GATA Haydarpasa Training Hospital, Istanbul, Turkey ; ⁶Infectious Disease and Clinical Microbiology, Gaziosmanpasa University Medicine Faculty, Tokat, Turkey ; ⁷Infectious Disease and Clinical Microbiology, Turkish Armed Forces Health Command Health and Veterinary Services, Ankara, Turkey ; ⁸Sarikamis Government Hospital, Kars, Turkey

ESCMID 2014

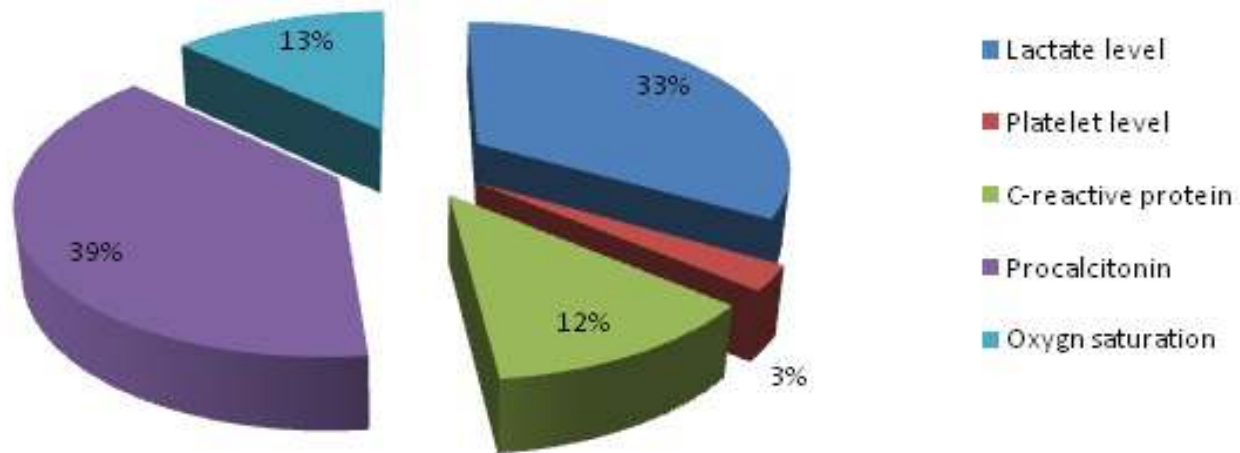


Figure 1. Q-Which parameter is more important than others in sepsis management?

- Ceftriaxone or other cephalosporins
- A carbapenem plus aminoglycoside
- Other
- Ceftriaxone plus vancomycine
- A carbapenem plus vancomycine

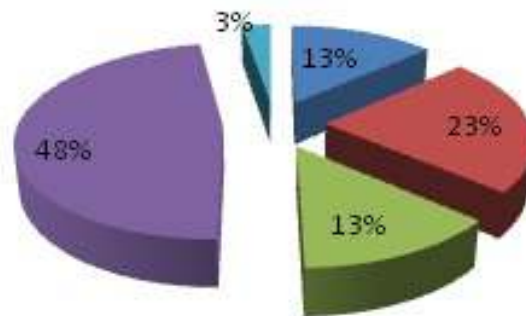


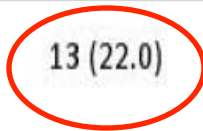

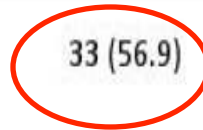





Fig 2. Q-Which antibiotics should be used in sepsis ?

Table 1. The rate of correct answers to sepsis bundle survey of different departments

	Infectious Diseases (59)	Emergency (46)	Internal Medicine (62)	Anesthesiology (56)
	n (%)	n (%)	n (%)	n (%)
Need for measuring lactate level	50 (81.4)	42 (92.3)	29 (46.8)	54 (96.4)
Take blood culture within 3 hours prior to antibiotics	58 (98)	 41 (89.1)	53 (85.5)	47 (83.9)
Administer crystalloid in hypotension (30 ml/kg)	 12 (20.3)	26 (56.5)	21 (33.9)	31 (55.4)
Lactate threshold in sepsis (>4mmol/L)	 13 (22.0)	 20 (43.5)	5 (8.2)	21 (37.5)
Target mean arterial pressure (≥65mmHg)	 33 (56.9)	 39 (84.8)	29 (46.8)	29 (51.8)
Target central venous pressure (8-12mmHg)	31 (52.5)	31 (67.4)	38 (61.3)	 41 (73.2)
Re-measure lactate if initial lactate was elevated	42 (71.2)	34 (73.9)	45 (72.6)	 46 (82.1)



WANTED

Sepsis

The largest
KILLER
Of children in the World