

The Effect of The Diagnostic Criteria on the Prognosis of Patients **Diagnosed With Sepsis at the Intensive Care Unit**

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Article info	Abstract	Research Article
Received: 08.06.2020 Received in revised form: 30.07.2020 Accepted: 10.08.2020 Available online: 05.09.2020	Sensis is a hard clinical condition to diagnose and tr	eat; but it occasionaly happens and it is an important life threatning health
Keywords	condition. New diagnostic criteriae has been release Sepsis Campaign diagnostic parameters and prognos and Research Hospital II. Anesthesiology and Rea	d in 2016. The aim of this study was to show the efficiency of Surviving tic factors. 100 patients in intensive care unit of Izmir Ataturk Education animation ICU were included. The data of the patients were recorded
Sepsis Intensive care unit CRP Procalcitonin	separately in both the survived (Group I) and decea each parameter with statistical significance. A total Among the SSC diagnostic criteria; respiratory rate hypoxemia, presence of acute oliguria, thrombocytop	iteria during the time interval of the study. Each parameter was assessed sed (Group II) patients. The specificity and sensitivity was calculated for of 100 patients were evaluated. The mortality of sepsis patients was 46%, , mental status, elevation of CRP and procalcitonin, presence of arterial benia, aPTT elevation, hypotension and lactate elevation were determined vities of CRP (82% and 95%, respectively) and of procalcitonin (53% and

INTRODUCTION

Sepsis is an important cause of morbidity and mortality in intensive care patients. Sepsis is a life threatning condition and defined as a response to infection due to unregulated/impaired host defence ¹⁻⁴.

that consensus meetings were held about diagnostic criteria and Surviving Sepsis Campaign should be carefully followed and treatment guidelines and their possible revisions. The new treatment should be initiated ^{5,6}. guideline was released in 2016^{-1} .

inflammatory markers did not have a place; but in 2017 are not specific to the disease. In addition, there have not been criteria they still have the importance.

Recently treatment protocols have been required due to an increment in hospitalizations due to sepsis and high morbidity and mortality rates.

Close monitoring in intensive care units enables early recognition of problems that may develop in these cases, and

contributes to the recovery process in patients. Early hemodynamic evaluation, assessment of physical findings, and abnormalities in parameters such as central venous pressure and urinary output help identification of persistent tissue hypoxia. In order to perform the targeted treatment The sepsis findings are nonspecific and because of immediately, the diagnostic criteria determined by the

82%, respectively) were high. The 2001 SSC diagnostic criteria are useful in determining mortality in patients with sepsis in intensive care unit. Nevertheless, pre-screening of high-risk patients, avoidance of unnecessary admissions, and creating

guidelines for approaching the patients with this disease will reduce mortality and morbidity rates.

There are a large number of diagnostic criteria, which According to the new sepsis criteria in 2016; are recognized to determine sepsis, but these diagnostic criteria a sufficient number of studies defining the superiority of any of these diagnostic criteria in terms of diagnosis, treatment and prognosis. The purpose of this study was to compare the roles of the 2001 diagnostic parameters in predicting mortality and to establish prognostic factors for a poor prognosis in patients admitted to our intensive care unit.

MATERIALS and METHODS

Ethical approval

The data in this study was obtained retrospectively from the hospital automation system and because of that no ethical approval was taken

In our study, following the approval by the local ethic committee of Izmir Ataturk Training and Research Hospital, the data of 100 patients, who were monitored between 01.September.2009 and 01.October.2010 with the diagnosis of sepsis at the 2nd anesthesiology and reanimation intensive care unit of Izmir Ataturk Research and Training Hospital, were recorded according to the diagnostic criteria developed by the 2001 Surviving Sepsis Campaign (SSC) and were evaluated mortality rate was calculated as 46%. In Group II patients, the retrospectively.

These diagnostic criteria are listed below (5-9):

Centigrade, b) Heart rate > 90/min, c) Tachypnea, respiratory rate > 22/dk, d) Altered mental status (GCS < 11), e) The recommendations were compared, it was determined that presence of Edema or positive fluid balance (> 20 mL/kg over among the global variables, the respiratory rate and GCS 24 hours), f) Hyperglycemia (glucose level > 120)

2. Inflammatory variables: a) Leukocytosis (WBC > 12,000), b) Leukopenia (WBC < 4000), c) elevation of CRP, d) Elevation of Procalcitonin

3. Hemodynamic variables: a) Hypotension (systolic blood pressure 90 mmHg, MAP < 70 mmHg or normal values less than 2 SD below normal for age)

4. Organ dysfunction: a) Arterial hypoxia (PaO2 / FiO2 > 300), b) The presence of acute oliguria (urine output < 0.5 ml/ kg/hour), c) Creatinine increase, d)Coagulation abnormalities (INR > 1.5 or aPTT > 60), e)Ileus, f) Thrombocytopenia (< 100,000/mm3), g)Hyperbilirubinemia (total bilirubin > 4 mg/dl)

5. *Tissue perfusion*: a) Hyperlactatemia (> 1mmol/L)

After the evaluation of demographic data, After the evaluation of demographic data, all parameters on the record were separated into 2 groups; one group consisting of discharged patients (Group I), the other group consisting of decedents (Group II). The statistical significance of each parameter was evaluated and the specificity and sensitivity of each parameter with a statistical significance were calculated.

Exclusion criteria: Patients below 18 years of age, pregnant patients with sepsis, patients with a previous established diagnosis of chronic renal failure and of diabetes were excluded from the study.

SPSS for Windows 17.0 program was used for statistical analysis. Besides the descriptive statistical methods, Chi Square test for the qualitative data, and Independent Samples T-Test for the quantitative data were used in comparisons. The parameters demonstrating statistical significance between the groups were evaluated by logistic regression analysis. The results within the 95% confidence interval and a value of p < 0.05 were considered statistically significant.

RESULTS

The study data demonstrated that 54 patients were discharged from the intensive care unit while 46 patients died. The age, the APACHE II and SOFA scores at the time of diagnosis were statistically significant compared to those in Group I 1. General criteria: a) Fever > 38 or <36 degrees patients (Table 1). When the diagnostic criteria for sepsis according to the Surviving Sepsis Campaign 2001 values (Table 2); among the inflammatory variables, procalcitonin and CRP values (Table 3); among the organ dysfunction variables, arterial hypoxemia; acute oliguria, and the platelet count (Table 4); in coagulation abnormalities, aPTT (Table 5); and hemodynamic and tissue perfusion variables (Table 6) were detected as the criteria with a prognostic value in Group II patients.

> The specificity and the sensitivity of the 11 diagnostic criteria, which were reported by the Surviving Sepsis Campaign in 2001, and which would have a potential to demonstrate a prognostic quality are demonstrated in Table 7. Of the diagnostic criteria, only two of them, namely procalcitonin and CRP levels were determined to have high sensitivity and specificity.

Table 1. Demographic data, APACHE II and SOFA scores

	Grup 1,(n=54)	Grup 2,(n=46)	Р
Gender (Female/Male)	18/36	25/21	>0.05
Age	64.95±13.33	55.94±18.09	=0.007*
APACHE II	22.86±8.78	27.25±5.75	=0.01*
SOFA	6.19±2.02	8.82±3.05	=0.05*

*, p<0.05

Table 2. General variables of the diagnostic criteria for sepsis

Pathology	Group 1, n (%)	Group 2, n (%)	Р
Fever	29 (53.7)	25 (52.1)	< 0.05
Heart rate	40 (74.1)	40 (86.9)	>0.05
Respiratory rate	24 (44.4)	30 (65.2)	=0.04*
Glasgow Coma Scale	7 (13)	18 (39.1)	=0.004*
Positive fluid balance	8 (14.8)	10 (21.7)	>0.05
Hyperglycemia	10 (18.5)	10 (21.7)	>0.05
*, p<0.05			

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Table 3. Inflammatory variables of the diagnostic criteria for sepsis

Pathology	Group 1,n (%)	Group 2,n (%)	Р
Leukocyte count	46 (85.2)	35 (76.1)	>0.05
Procoalsitonin	29 (53.7)	38 (82.6)	=0.002*
CRP	44 (81.5)	44 (95.6)	=0.025*

*, p<0.05

Table 4. Organ dysfunction variables of the diagnostic criteria for sepsis

Pathology	Group 1,n (%)	Group 2,n (%)	Р
Arterial hypoxia	25 (46.3)	32 (71.1)	=0.02*
Acute oliguria	6 (11.1)	14 (31.1)	=0.02*
Kreatinin	15 (27.8)	21 (46.7)	>0.05
Ileus	12 (22.2)	26 (35.6)	>0.05
Thrombocyte count	5 (9.3)	12 (26.7)	=0.04*
Biluribin level	14 (25.9)	19 (42.2)	< 0.05

*, p<0.05

Table 5. Coagulation abnormalities of the diagnostic criteria for sepsis

Pathology	Group 1,n (%)	Group 2,n (%)	Р
INR	12 (22.2)	18 (39.1)	< 0.05
aPTT	3 (5.6)	11 (23.9)	=0.01*
*, p<0.05			

Table 6. Hemodynamic and tissue perfusion variables of the diagnostic criteria for sepsis

Pathology	Group 1,n (%)	Group 2,n (%)	Р
SAB	14 (25.9)	28 (62.2)	=0.001*
OAB	13 (24.1)	27 (58.6)	=0.001*
Lactate level	8 (14.8)	27 (58.6)	<0.000*

*, p<0.05

 Table 7. The sensitivity and specificity rates of the diagnostic criteria with statistical significance

	Sensitivity (%)	Specifity (%)
Respiratory rate	65	55
Glasgow coma scale	39	12
Procalsitonin	82	53
CRP	95	81
Arterial hypoxia	69	46
Acute oliguria	45	11
Thrombocyte count	26	1
aPTT	24	0.05
SAB	60	24
OAB	58	17
Lactate level	58	17

DISCUSSION

Sepsis is a hard clinical condition to diagnose and treat; but it occasionaly happens and an important life threatning health condition ^{6,10,11,12}. According to the new sepsis criteria in 2016; inflammatory markers did not have a place; but in 2017 criteria they still have the importance ¹⁻⁴. The annual rate of sepsis is 50-95 cases per 100,000, and this figure shows an increase by 9% every year ⁶. In the international literature, the mortality of severe sepsis has been reported as 30-50%. The mortality risk in sepsis patients is affected by disease severity, age, and underlying diseases ⁶. Poeze et al. ¹³ conducted a study by addressing questions on a phone call to randomly selected 10058 physicians from Europe and U.S. The interviewed

physicians were either intensive care unit physicians or physicians, who spent 50% of their duration of working time in treating intensive care unit patients. The authors reported that 67% of the physicians thought that there are deficiencies in applying a common definition of sepsis, and 83% of them thought that sepsis cases were often missed as they could be confounded by symptoms of many diseases. Indicating factors of poor prognosis of sepsis especially will be more helpful in terms of treatment and monitoring ^{14,15}. The performance of these criteria, namely biomarkers, are measured by sensitivity, which is the ability to detect the patients with the correct diagnosis; and by specificity, which is the ability to recognize the patients without a correct diagnosis ¹⁶. But there are some problems in the biomarkers and in the criteria for sepsis in terms of sensitivity and specificity ^{17,18}. Therefore, we aimed to compare the effects of diagnostic parameters published in 2001 on the mortality estimates of the sepsis patients admitted to our intensive care unit; and to identify the poor prognostic factors in order to contribute to diminish the problems of making an initial diagnosis and to the decline of mortality rates.

In our study, 100 patients admitted to the reanimation clinic of the anesthesiology and reanimation department with the diagnosis of sepsis were evaluated. Of these 100 patients, 54 of them were discharged from the intensive care unit, while the 46 of them died, demonstrating a mortality rate of 46% in accordance with the literature.

Sepsis, is common in advanced ages, especially above the age of 50 $^{10-12}$. It is more common in men than women. The reason for this difference is unknown, however, it is suggested that men are more prone to trauma, and to undergo surgery. In our study too, sepsis was detected in male patients at an advanced age.

CRP levels can be elevated in diseases other than sepsis (trauma, burns, surgery, infarction) ^{19,20}. In spite of this fact, elevations in CRP levels were demonstrated to be a poor prognostic factor by the study by Lobo et al. ¹⁹ and by other studies investigating the effect of this protein ²⁰⁻²². Our study results concluded that increased CRP was a significant criterion for the diagnosis and prognosis. Inflammatory conditions, localized bacterial infections, and viral infections do not trigger the increase of procalcitonin extensively, whereas procalcitonin ratio is elevated in systemic bacterial infections seriously ²³⁻²⁵. As the sensitivity and specificity of Pro-Cal are reported ²³⁻²⁷ to specific than CRP in many studies, debate continues ²⁸. Simon suggestion. et al.²⁴ reports that it is a more successful method in defining significant effect of Pro-Cal on mortality is an indicator that found to be useful and significant on prognosis. this protein may be effective in the prognosis of sepsis among with high levels of specificity and sensitivity.

 $(\geq 37.8^{\circ}C)$ and heart rates of ≥ 90 are significant to diagnose prognosis of sepsis. bacteremia, they cannot be established as data influencing the prognosis of sepsis directly. Our study results support this the diagnostic criteria for sepsis published in 2001 respiratory approach as no significant results were obtained for fever and rate, GCS, CRP, procalcitonin levels, systolic arterial pressure, heart rate. Sensitivity and specificity rates of fever were mean arterial pressure, arterial hypoxemia, oliguria, calculated as 53% and 46%, respectively, and did not coagulation disorders (APTT), platelet count, MVS, age, and constitute a confidence level.

Our study results demonstrate that the change in fluid fever, leukocyte count, balance, increase in glucose, creatinine, presence of a hyperglycemia, creatinine levels, ileus, hyperbilirubinemia, and coagulation disorder (INR), bowel movements, total bilirubin INR values have no effect on the prognosis. To be change, and the presence of comorbid diseases are not acknowledged of the poor prognostic factors is important in significant in the prognosis of sepsis. Although the presence of admissions to intensive care unit and during monitoring the a comorbid disease has been demonstrated as a criterion patients. It is suggested that future guidelines, to be developed accelerating or contributing to mortality, it is not an effective for the treatment management of high risk patients by taking all data. This is interpreted as the result of excluding the patients of these parameters/variables into consideration, will decrease with diabetes and chronic renal failure, which might have a the mortality and morbidity rates. By means of the treatment negative effect on the prognosis, and as the non-significant guidelines, the redundant admissions of the low risk patients to effects of comorbidities such as hypertension and chronic the intensive care unit, or the delayed intensive care obstructive lung disease on the prognosis of sepsis.

The factors demonstrated to trigger sepsis include endothelial activation independent of global tissue hypoxia, impaired balance in the coagulation mechanisms, increased vascular permeability, and impaired vascular tone. Respiratory support to the patients in the intensive care units is a factor in obtaining favorable outcomes in the treatment 13,30 . Among the 1. criteria investigated by our study, coagulation disorder (aPTT), arterial hypoxemia, and changes in respiratory rate are demonstrated to be the potential significant prognostic 2. indicators.

As the study results of Meyancı et al. ⁵ demonstrate that the changes in urinary output are effective in detecting persistent tissue hypoxia, investigation of this data in sepsis is

be 65-97% and 48-94% respectively in the study reports, it is suggested to have a potential in providing significant determined that these values are not sufficient to make a information. Oliguria, which is among the criteria investigated diagnosis. Although procalcitonin is recognized as being more in our study revealed significant results thereby supporting this

In parallel to the results of the studies by Nguyen et al. the severity of sepsis and its prognosis. A statistically and H-Michael et al ^{31,32}, the change of lactate levels has been

By the clinical procedures and the studies conducted, the specified criteria ^{26,27,29}. The results of our study, too, it is acknowledged that the thrombocyte count and thrombocyte demonstrates that CRP and procalcitonin are the two criteria supply carry the significance with them in the disease prognosis ^{5,6,33}. Therefore, the changes in thrombocyte levels as Although Jaimes et al.²⁹ determines that high fever a criterion is determined to be a significant indicator in the

> As a result of our study, it is concluded that, among lactate levels influence the prognosis. It is determined that heart rate. fluid balance. monitorization of the critical patients can be prevented.

Conflict of interest

The authors declare that they have no conflict of interest.

REFERENCES

- Singer M, Deutschman CS, Seymour CW et al. The Third International Consensus Definitions for Sepsis and Septic Shock (Sepsis-3). JAMA 2016; 315(8):801-10.,
- Shankar-Hari M, Phillips GS, Levy ML et al. Developing a new definition and assessing new clinical criteria for septic shock: for the Third International Consensus Definitions for Sepsis and Septic Shock (Sepsis-3). JAMA .2016; 315(8):775-87.

- 3. Seymour CW, Liu VX, Iwashyna TJ et al. Assessment of clinical 18. Calandra T, Cohen J: The international sepsis forum consensus criteria for sepsis: for the Third International Consensus Definitions for Sepsis and Septic Shock (Sepsis-3). JAMA 2016; 315(8):762-74.)
- Rhodes A, Evans LE, Alhazzani W, Levy MM, Antonelli M, Ferrer 4. R et al: Surviving Sepsis Campaign: International Guidelines for Management of Sepsis and Septic Shock: 2016. Crit Care Med. 2017 Mar;45(3):486-552.
- 5. Meyancı G. Early Targeted Therapy in Severe Sepsis and Septic Shock. Sepsis Symposium Series in the Light of Current Information May 2006; 51: 45-9.
- Dellinger RP, Carlet JM, Masur H, et al. Surviving Sepsis Campaign 6. guidelines for management of severe sepsis and septic shock. Crit Care Med .2004; 32: 858-73.
- J, Opal SM, Vincent JL, Ramsay G: 2001 SCCM/ESICM/ACCP/ATS/ 22. Seller Perez G, Herrera-Gutierrez ME, Lebron- Gallardo M, 7. SIS International Sepsis Definitions Conference. Intensive Care Med. 2003, 29: 530-38.
- 8. Levy MM, Fink MP, Marshall JC, Abraham E, Angus D, Cook D, Cohen J, Opal SM, Vincent JL, Ramsay G: 2001 SCCM/ESICM/ 23. Carrol ED, Thomson APJ, Hart CA. Procalcitonin as a marker of ACCP/ATS/SIS International Sepsis Definitions Conference. Crit Care Med 2003; 31: 1250-56.
- Calandra T, Cohen J: The international sepsis forum consensus 9. conference on definitions of infection in the intensive care unit. Crit Care Med 2005; 33: 1538-15
- 10. Aygün G. Sepsis Diagnosis Sepsis Symposium Series in the Light 25. Ugarte H, Silva E, Mercan D, de Mendonca A, Vincent JL: of Current Information May 2006; 51: 51-60.
- 11. Mitaka C. Clinical laboratory differentiation of infectious versus non-infectious systemic inflammatory response syndrome. 26. Redl H, Schlag G, Togel E, Assicot M, Bohuon C: Procalcitonin Clinica Chimica Acta 2005; 35: 17-29.
- 12. Pierrakos C, Vincent JL, Sepsis biomarkers: a review. Critical Care 2010; 14: 15, 18.
- 13. Poeze M, Ramsay G, Gerlach H Rubulotta, Lewy M : An 27. Becker KL, Snider R, Nylen ES: Procalcitonin assay in systemic Internati- onal Sepsis Survey ; A Study of Doctors Knowledge and Perception about Sepsis . Critical Care 2004; 435-6
- 14. Rhodes A, Bennett D. Early goal-directed therapy: An 28. Pierrakos C, Vincent JL, Sepsis biomarkers: a review. Critical Care evidence-based review. Crit Care Med. 2004; 32; 448-50
- 15. Rivers E, Nyugen B, Haustad S et al. Early goal of directed 29. Jaimes F, Arango C, Ruiz G, et al. Predicting bacteremia at the the- raphy in the treatment of severe sepsis and septic shock. N Eng J Med. 2001; 53; 1368-77
- 16. Marshall JC, Reinhart K, Biomarkers of sepsis. Crit Care Med 2009; 37: 2290-98

Opal SM, Vincent JL, Ramsay G: 2001 SCCM/ESICM/ACCP/ATS/ 31. Nguyen HB, Rivers EP, Knoblich BP, Jacobsen G, Muzzin A, SIS International Sepsis Definitions Conference. Intensive Care Med. 2003, 29: 530-38.

- conference on definitions of infection in the intensive care unit. Crit Care Med 2005, 33: 1538-48.
- 19. Lobo SM, Lobo FR, Bota DP, Lopes-Ferreira F, Soliman HM, Melot C, Vincent JL: C-reactive protein levels correlate with mortality and organ failure in critically ill patients. Chest 2003, 123: 2043-49
- 20. Schmit X, Vincent JL: The time course of blood C-reactive protein concentrations in relation to the response to initial antimicrobial therapy in patients with sepsis. Infection 2008, 36:213-19.
- 21. Couto RC, Barbosa JA, Pedrosa TM, Biscione FM: C-reactive pro- tein-guided approach may shorten length of antimicrobial treat- ment of cultureproven late-onset sepsis: an intervention study. Braz J Infect Dis 2007, 11: 240-45
- Toro- Peinado I, Martin Hita L, Porras Ballesteros JA; Serum CRP as a marker of outcome and infection in critical care patients. Med Clin Care 2005; 125: 761-65
- sepsis. Int J Antimicrobial Agents 2002; 20: 1-9.
- 24. Simon L, Gauvin F, Amre DK, Saint-Louis P, Lacroix J. Serum pro- calcitonin and C-reactive protein levels as markers of bacterial infection: a systematic review and meta-analysis. Clin Infect Dis 2004; 39: 206-17
- Procal- citonin used as a marker of infection in the intensive care unit. Crit Care Med 1999, 27: 498-504.
- release patterns in a baboon model of trauma and sepsis: relationship to cytokines and neopterin. Crit Care Med 2000, 28: 3659-63.
- inflammation, infection, and sepsis: clinical utility and limitations. Crit Care Med 2008, 36: 941-52.
- 2010, 14: 15, 18.
- bedside. Clin Infect Dis 2004; 38:357-62.
- 30. Rivers E ,Nyugen B, Haustad S et al. Early goal of directed the- raphy in the treatment of severe sepsis and septic shock. NEng J Med 2001; 53; 1368-77
- Ress- ler JA, Tomlanovich MC: Early lactate clearance is associated with improved outcome in severe sepsis and septic shock. Crit Care Med 2004, 32:1637-42.

- 32. Haji-Michael PG, Ladriere L, Sener A, Vincent JL, Malaisse WJ: Leukocyte glycolysis and lactate output in animal sepsis and ex vivo human blood. *Metabolism* 1999, 48:779-85. Haji-Michael PG, Ladriere L, Sener A, Vincent JL, Malaisse WJ: Leukocyte glycolysis and lactate output in animal sepsis and ex vivo human blood. Metabolism 1999, 48:779-85.
- Ürkmez S, Survival in Sepsis Campaign. İ.Ü.Cerrahpaşa Faculty of Medicine, Department of Anesthesiology, Sadi Sun Intensive Care Unit: *Steteskop Net* 2006;6: 27-28