Procalcitonin (ProCT): A Reliable Marker of Sepsis. A Single Centre Experience

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Abstract: Objective: Sepsis occurs in hematological disorders due to neutropenia, clinical signs and symptoms are not specific for its diagnosis. Early biomarkers are potentially helpful for the diagnosis in these patients. This study was done to assess the utility of Procalcitonin (ProCT) as a marker of sepsis in patients with hematological disorders.

Methods: A cross sectional study was conducted in the section of clinical chemistry, department of laboratory, National institute of Blood disease and Bone marrow transplantation (NIBD) Hospital in Karachi Pakistan. Total 366 patients with age groups of 15 to 60 years with suspected sepsis undergone ProCT test were enrolled in the study, their blood cultures results and biomarkers including; White blood cell (WBC) count, C-reactive protein (CRP) and Absolute Neutrophil count (ANC) were also recorded. Sepsis was labeled as per surviving sepsis campaign (SSG) international guidelines for management of severe sepsis and Septic Shock 2012 criteria.

Results: Biomarkers of culture positive patients were compared with culture negative patients. The statistically significant difference was noted in ProCT (p value < 0.001) and CRP (p value < 0.003). However, no significant differences were noted in WBC count (p value > 0.05) and ANC count (p value > 0.05). ProCT had outstanding diagnostic accuracy with area under receiver operator characteristic curve (AUC) was 0.95 ; (95%CI0.89 - 1.00), that for CRP was modest (AUC 0.74; 95%CI 0.61-0.87), while that for WBC and ANC were poor with AUC 0.43; (95%CI 0.31-0.58), AUC 0.44(95%CI 0.28-0.54) respectively.

Conclusion: ProCT alone showed the excellent diagnostic power on ROC analysis which proves it as a stronger diagnostic biomarker for sepsis in hematological disorders.

Keywords: Procalcitonin (ProCT), Suspected sepsis, Hematological disorders, White blood cell (WBC), C-reactive protein (CRP), Absolute neutrophil count (ANC), Neutropenia.

INTRODUCTION

Sepsis is a massive immune response to bacterial infection, which affects many organs and without prompt management-can lead to shock and multisystem organ failure [1]. It is the second most common cause of death after myocardial infarction in patients admitted to intensive care units [2]. Clinical, laboratory and microbiologic strategies are used for its diagnosis. Laboratory tests ranging from white blood cell count (WBC), neutrophil count and erythrocyte sedimentation rate (ESR), to more complex biomarkers such as blood cultures, Procalcitonin (ProCT), cytokine levels and circulating cell-free DNA(cfDNA) are used. The common approach to detect sepsis is by blood cultures and they are considered as gold standard, but it has certain limitations as they are positive in up to 30% of patients and takes 2 to 3 days for the results to be available [3, 4]. Several studies have reported negative culture results even in severe sepsis, resulting in difficulty in diagnosing such cases.

The diagnosis of infection is complex and sometimes impossible because of limited sign and symptoms in patients receiving intensive chemotherapy for hematopoietic stem cell transplant (HSCT) which leads to long-term granulocytopenia. Fever is the only sign of infection in these patients [5]. Identifying the origin of a fever is very essential to start appropriate therapy. The working group of the German Society of Hematology and Oncology suggest that in neutropenic patients, the white blood cell count should not be used in criterion to define sepsis. The Third International Consensus Definitions for Sepsis and Septic Shock stated that sepsis is a clinical diagnosis with no gold-standard diagnostic test [6].

For the above mentioned reasons, a reliable biomarker is needed to identify sepsis, that must be sensitive, specific, predict early diagnosis and easy to measure and useful in neutropenic patients as well. Since recent decades ProCT has emerged as a highly sensitive biomarker for the diagnosis of sepsis. Normal serum ProCT levels in the blood are very low. However its concentration instantaneously elevates in 6 to 12 hours of infection and after treatment, its levels declines and with undetectable levels in patients with complete recovery [7, 8]. Its levels are proportional to the severity of infection, higher levels associated with more severe disease and vice versa. In 2001 International Sepsis Definitions Conference added ProCT and C-reactive protein to the new diagnostic criteria of sepsis [9]. Centers for Medicare & Medicaid Services (CMS) regulations have stated that ProCT is a

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reliable and accurate biomarker for sepsis diagnosis [10].

Diagnostic accuracy of ProCT in patients with neutropenia is not similar in studies [7]. Prior systematic analysis and meta-analyses were based mainly on pediatric population [11-13], and literature on adult population is not sufficient. Keeping in mind all these things, we planned our study to define the clinical significance of serum ProCT in Septic patients.

MATERIALS AND METHODS

Patients’ Recruitment

This was a cross sectional study, conducted in a section of clinical chemistry, Department of Laboratory, National Institute of blood disease and bone marrow transplantation (NIBD) Hospital, Karachi Pakistan from 1st December 2017 to 1st June 2018. The study was conducted after approval of the Institutional Review Board (IRB)/Ethical Committee of Hospital with Protocol no NIBD/RD-181/2018. Patient demographics and relevant biochemical data were recorded on a structured proforma.

Inclusion Criterion

Patients with age 15 to 60 years undergoing ProCT test with suspected sepsis during the study period were enrolled in the study. Sepsis was defined as per Surviving Sepsis Campaign (SSG) International Guidelines for Management of Severe Sepsis and Septic Shock 2012 criteria defined sepsis as the presence (probable or documented) of infection together with systemic manifestations of infection, temperature >38°C or <36°C, heart rate more than 90 beats per minute, respiratory rate of more than 20 breaths per minute and white blood cell count >12 000/µL or <4000/µL or >10% immature (band) forms. Traditionally, more than two SIRS criteria were felt to represent patients at risk of or suffering from sepsis [12, 14].

Exclusion Criterion

Pregnant females, patients with sepsis and already started on antibiotic treatment, the death of patients due to sepsis and unavailability of blood culture reports were excluded.

Biomarkers Analysis

Three to five milliliter of blood was collected in yellow top (gel tube) and ProCT analyzed by electrochemiluminescence technique on Roche Cobas e-411 analyzer, cut off used as per Consensus conference of the American College of Chest Physicians/Society of Critical Care Medicine. ProCT levels < 0.5ng/ml considered as low risk for sepsis, ProCT levels ≥ 0.5ng/ml but <2ng/ml considered as moderate risk for sepsis, ProCT levels ≥2 and <10ng/ml considered as high risk for sepsis, ProCT levels ≥10ng/ml considered as severe sepsis or septic shock [14-16]. CRP was analyze by Particle enhanced immunoturbidimetric assay on Roche Cobas c-111 analyzer, cut off used CRP<10mg/L labeled as no infection while CRP>10mg/L labeled as infection. WBC and Neutrophil count was analyzed on XN-1000 Sysmex Hematology analyzer by flow cytometry technique. The cut off used for ANC was <2x109/L considered as neutropenia, 2-7x109/L considered normal ANC and >7x109/L considered as neutrophilia. For Internal quality control; two levels of manufacturer provided controls (low and high) were run of ProCT, CRP, WBC, and Neutrophil count (ANC) laboratory also participated in RIQAS (Randox International quality assurance scheme) as External quality assurance program for proficiency testing.

ANALYSIS

Statistical software SPSS version 19 was used for analysis of data. Frequency and percentages were calculated for gender, while mean and standard deviation were calculated for age. Median and interquartile (IQR) were calculated for ProCT, CRP, WBC, and ANC count. The ProCT, CRP, WBC and ANC levels were compared between culture positive and cultures negative patients by using Mann-Whitney U test. Kappa statistic was used to find the agreement between blood culture and Procalcitonin. Diagnostic accuracy of ProCT, CRP, WBC and ANC to predict sepsis was calculated against gold standard as blood culture. Receiver operator curve and area under the curve was derived. A p-value of <0.05 was considered as significant.

RESULTS

Patients undertaken serum ProCT test with suspected sepsis during the period 1st December 2017 to 1st June 2018 were included in the study. Total 366 subjects were recruited, out of which 296 were excluded due to unavailability of blood cultures results. Total 70 subjects were included in final analysis with mean age of 36.5±16.3years, 57.1% (n=40) were males and 42.8% (n=30) were females. The median serum CRP level was 15.4mg/L (IQR: 3.2-84.3), median ProCT was 0.40ng/ml (IQR: 0.121-1.2), median WBC was 2.4x109/L (IQR: 0.79-6.65) and mean ANC was 1.5x109/L (IQR: 0.151-3.94). The 57.1% (n=40) patients had ANC count <2x109/L (Neutropenic), while 31.4% (n=22) had normal ANC and 11.4% (n=8) patients had ANC more than 7x109/L (Neutrophilic).

There were (n=24) 34.2%culture positive, while (n=46) 65.7% were cultures negative patients. Out of 24 cultures positive patients, ProCT correctly identified 22 (91.6%) patients that levels more than 0.5ng/ml. Out of 46cultures negative patients ProCT correctly identified 37(80.4%) that had ProCT levels less than 0.5ng/ml as shown in Table 1. ProCT, CRP, WBC and ANC were compared between positive and negative blood culture groups, a significant difference was seen with ProCT (p-value <0.001), and CRP (p-value 0.003), whereas no significant differences were seen in WBC count (p-value 0.16) and ANC (p-value 0.23) as shown in Table 2. On ROC analysis, the AUC of serum
ProCT was 0.93 (95% CI 0.87-0.99), CRP was 0.71 (95% CI 0.58-0.85), WBC0.39 (95% CI 0.25-0.54) and for ANC was 0.41 (95% CI 0.26-0.56) as shown in Fig. (1). At a cut-off of 0.5ng/ml ProCT had sensitivity of 95.5% specificity of 78.2%, PPV of 69.7% and NPV of 97.3%, the positive likelihood ratio for Procalcitonin 4.5 (95% CI 2.58-7.85) while the negative likelihood ratio 0.06 (95% CI 0.01-0.80) with accuracy of 84.29(95% CI 73.62-91.89). There were positive agreement between blood cultures and Procalcitonin with kappa value of 0.676 with p-value of <0.001.

Table 1. Accuracy of Procalcitonin (ProCT) at Cut off 0.5ng/ml.

<table>
<thead>
<tr>
<th>Test</th>
<th>Blood culture positive (n=24)</th>
<th>Blood culture negative (n=46)</th>
<th>Total (n=70)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Procalcitonin Positive</td>
<td>22</td>
<td>10</td>
<td>33</td>
</tr>
<tr>
<td>Procalcitonin Negative</td>
<td>01</td>
<td>37</td>
<td>37</td>
</tr>
</tbody>
</table>

Table 2. Sepsis Markers of Patients with Positive and Negative Blood Cultures.

<table>
<thead>
<tr>
<th>Sepsis Markers</th>
<th>Blood culture positive Median(IQR)</th>
<th>Blood culture negative Median(IQR)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>ProCT (ng/ml)</td>
<td>3.1 [0.94-20]</td>
<td>0.2 [0.09-0.44]</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>CRP (mg/L)</td>
<td>79.9 [15-171]</td>
<td>7.2 [3.0-41.6]</td>
<td>0.002</td>
</tr>
<tr>
<td>WBC (x10^9/L)</td>
<td>1.34 [0.38-5.0]</td>
<td>2.4 [1.1-7.0]</td>
<td>0.16</td>
</tr>
<tr>
<td>ANC (x10^7/L)</td>
<td>0.34 [0.0-3.8]</td>
<td>1.7 [0.24-4.2]</td>
<td>0.46</td>
</tr>
</tbody>
</table>

DISCUSSION

ProCT is 116-amino acid polypeptide prohormone of calcitonin synthesized by the C-cells of the thyroid gland, and in small amount by neuroendocrine tissue, lungs and intestines. Normally ProCT levels in the blood are very low; its production is stimulated by inflammatory cytokines and bacterial endotoxins causing increased levels in the blood. It has superior biokinetics, characterized by increased levels earlier in bacterial infection or sepsis and decrease faster when infection is controlled.

In this study, the diagnostic utility of serum ProCT for the diagnosis of sepsis in hematological disorders was evaluated. All subjects with suspected sepsis with hematological disorders undergoing serum ProCT test were included in the study. It was observed that median serum ProCT level, 3.1ng/ml [IQR 0.94-20] was significantly higher in blood culture positive patients in comparison to blood culture negative patients, 0.2ng/ml [IQR 0.09-0.44] (p-value <0.001).

A study performed by M. Assicotand his colleagues in 1993 found very high concentrations of serum ProCT at diagnosis in severe bacterial infections patients (range 6-53 ng/mL) [17].

We used a cut off of serum ProCT 0.5ng/ml for sepsis diagnosis. Similar Cut-offs used by Vincenzi I. and his colleagues in 2016 and Romano his colleagues in 2019 [18, 19]. We reported a sensitivity of 96% and specificity of 22% at a cutoff point of 0.5 ng/ml sepsis diagnoses. However, by lowering the cutoff point to 0.1ng/mL led to increased sensitivity and specificity, 100% and 70% respectively. Romano and his colleagues in 2018 reported similar sensitivity and specificity of 100 and 22% respectively at a similar cut-off of 0.5ng/ml [19].

A systematic review of randomized controlled trials done by Schuetz P and his colleagues in 2011 stated that for optimal performance, serum ProCT cut-offs must be used according to patient risk level and clinical setting. Low-risk patients, typically those with respiratory tract infections presenting to an emergency department (ED), a serum ProCT cut-off of ≤0.25 ng/mL or 0.1 ng/mL should be used. While in high-risk patients, typically those admitted to the intensive care unit (ICU) serum ProCT level of 0.5 ng/mL or ≥0.26 ng/mL should be used [6].

Normally, serum ProCT levels are very low <0.01 ng/mL. In sepsis itsevels increases significantly, sometimes to more than several hundred nanograms per milliliter. Paul E. Marik in 2014 stated that ProCT level >0.5 ng/ml is highly suggestive of a bacterial infection whereas a level <0.1 ng/ml less likely to be bacterial infection [4].

A multicenter randomized trial done by De Jong (2016) and his colleagues on Stop antibiotics on procalcitonin guidance Study (SAPS), in patients with suspected infection or proven infection admitted to the ICU. They recommended that...
clinicians should stop antibiotics if ProCT is ≤0.5 μg/L or if it decreased by ≥80% of the peak value [7]. Riedel in 2011 stated that ProCT is extremely sensitive biomarker for sepsis diagnosis or to rule-out the bacterial sepsis, with a high negative predictive value of more than 95% [20].

The study has many limitations as 1) it was conducted at single institute with a small sample size 2) Serum ProCT mislabeled 14.2 % (n=10) out of 70 patients as septic and their levels were between 0.5 and 2 ng/ml. This could probably be due to low levels of bacteremia and bacteria failed to grow in culture, anaerobic bacterial infection or patients had some other cause of infection. One patient was missed by serum ProCT; it may be due to skin flora contaminant or maybe empirically antibiotic was started prior to blood culture, which resulted in a reduction of the serum ProCT values. 3) Most of the patients were post chemotherapy and post Bone Marrow Transplant (BMT) patients so their ANC and WBC did not rise significantly.

The strength of the study was a strict inclusion criterion of patients with suspected sepsis and one type of population included in study.

CONCLUSION
In conclusion, ProCT is a reliable and specific sepsis biomarker. It can be used as a single reliable test and it has about the same accuracy as blood cultures and more accurate than CRP, ANC, and WBC for sepsis diagnosis in patients with hematological disorders.

STUDY APPROVAL
Study commenced after the approval of Institutional Review Board (IRB)/Ethical Committee of NIBD Hospital with protocol no. NIBD/IR-181/2018.

DATA AVAILABILITY
Data is confidential.

FUNDING SOURCE
No.

AUTHORS’ CONTRIBUTIONS
1. Shabnam Dildar (SD) planned the study, collected and analyzed the data, and wrote the manuscript
2. Jawad Hassan (JH) critically reviewed the manuscript for intellectual content.
3. Tahir Sultan Shamshi (TS) critically reviewed the manuscript for intellectual content.

ABBREVIATIONS
AUC: Area under Curve.
cfDNA: Circulating Cell-Free DNA.
CI: Confidence Interval.
CRP: C-Reactive Protein.

ProCT: Procalcitonin.
SIRS: Systemic Inflammatory Response Syndrome.
BMT: Bone Marrow Transplant.
ANC: Absolute Neutrophil Count.

CONFLICT OF INTEREST
Declared none.

ACKNOWLEDGEMENTS
Declared none.

REFERENCES
Sepsis is a massive immune response to bacterial infection, which can lead to organ failure and can be life-threatening. The Surviving Sepsis Campaign (SSC) guidelines, published by the Society of Critical Care Medicine (SCCM) and the American Thoracic Society (ATS), define sepsis as a systemic inflammatory response syndrome (SIRS) associated with infection.

INTRODUCTION

Sepsis is defined as a syndrome characterized by systemic inflammation and organ dysfunction caused by an infection. The SSCC guidelines recommend early recognition and aggressive management of sepsis to improve patient outcomes. Procalcitonin (ProCT) is a biomarker that is widely used in the diagnosis and management of sepsis.


cell-free DNA (cfDNA) are used. The common approach in high-risk patients, typically those admitted to the intensive care unit (ICU), is to check for early signs of sepsis, such as fever, tachycardia, tachypnea, and hypotension.

Procalcitonin (ProCT) is a biomarker that is widely used in the diagnosis and management of sepsis. ProCT is a low-molecular-weight peptide that is produced by the thyroid gland and is involved in the regulation of the immune system. ProCT is released into the bloodstream in response to bacterial infection, and its levels increase in the blood in the early hours of infection.

The SSCC guidelines recommend that ProCT levels >0.5ng/ml is highly significant. ProCT levels <0.5ng/ml considered as low risk for sepsis, ProCT levels >0.5ng/ml are considered as an early sign of sepsis.

Early diagnosis of sepsis is crucial to prevent organ failure and mortality. ProCT is a promising biomarker for the diagnosis and management of sepsis, and it is widely used in clinical practice. However, ProCT levels can also be elevated in non-infectious conditions such as inflammation, trauma, and cancer.

In this study, the diagnostic utility of serum ProCT for early detection of sepsis was evaluated. All subjects with suspected sepsis with hematologic cultures included in the study. Total 366 subjects were recruited, out of which 296 were excluded due to unavailability of blood cultures, anaerobic bacterial infection or patients had some other cause of infection. One patient was missed by serum culture, anaerobic bacterial infection or patients had some other cause of infection. Most of the patients were post chemotherapy and post Bone marrow transplant (BMT).

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In conclusion, ProCT is a reliable and specific sepsis biomarker. It helps to diagnose early in bacterial infection or sepsis and decrease faster earlier in bacterial infection or sepsis and decrease faster. The SSCC guidelines recommend early recognition and aggressive management of sepsis to improve patient outcomes. ProCT is a promising biomarker for the diagnosis and management of sepsis, and it is widely used in clinical practice. However, ProCT levels can also be elevated in non-infectious conditions such as inflammation, trauma, and cancer.