Serum Procalcitonic (PCT) Versus Serum C - Reactive Protein (CRP) for Severity of Organ Dysfunction in Sepsis

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Abstract

Background: Intensive care units (ICUs), despite advances in critical care nursing, have frequent issues with early diagnosis and adequate treatment. Recently discovered world-class procalcitonin (PCT), a revolutionary laboratory marker, has been shown to be useful in this regard. The objective is to Comparison of concentrations of serum procalcitonin (PCT) and c-reactive protein (CRP) with a comparable level of organ malfunction during sepsis and evaluation of the interaction between serum PCT and CRP concentrations with different organ malfunction occurrence in sepsis. Design: It is a Hospital-Based Prospective study. Participants and Setting: Fifty people were admitted to the intensive care unit of Gandhi Medical College. **Subjects and Methods:** The extent of sepsis-related organ impairment was evaluated with the sequential organ failure assessment (SOFA) on day 1. Patients were identified by category 1(0-6), category 2(7-12), group 3(13-18), and group 4(19-24) in 4 separate classes with varying organ impairment seriousness of sepsis. Serum PCT and CRP concentrations have been measured. **Results:** The majority of the patients belonged to the age groups of 60-69 years (30%) and 50-59 years (22%) Majority of the patients belonged to the Sofa group 1 around 42% followed by sofa group 2 with 38%, sofa group 3 with 16% and the least belonged to the sofa group 4 with 4%. The mean PCT and CRP concentration in those who survived was 14.73 ng/ml and 149.916mg/L respectively and in those who died were 45.76 ng/ml (p-value <0.001) and 183.584 mg/L (p-value 0.172) respectively. The linear correlation between PCT plasma concentrations and the four groups was significantly stronger than with CRP. **Conclusion:** In SOFA and serum PCT, The level of organ dysfunction and complications in sepsis patients is closely related to serum CRP levels.

Keywords: SIRS (Systemic Inflammatory Response Syndrome), SOFA (Sequential Organ Failure Assessment), Sepsis, C - reactive protein (CRP), Procalcitonin (PCT)

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Introduction

A Greek phrase-putrid incorporates the name sepsis. Air touch and mortality were solely responsible for the putrefaction of a wound when the putrefaction cycle entered the blood (septicemia). PCT is not attributed to systemic bacterial diseases, respiratory illnesses, inflammatory and allergic conditions. It is not presently known if PCT is primarily affected by microbial inflammation of the extent of multiple organ dysfunctions due to systemic inflammatory reactions.

The criteria for the American College of Chest Physicists (ACCP) and the Society of Critical Care Medicines (SCCM),^[1] defines sepsis as the incidence of documented or alleged infectious systemic inflammatory reaction syndrome

with SIRS, [2,3] (systemic inflammatory response syndrome) identified by the existence of > 2:

- 1) Heart rate >90/min
- 2) Respiratory rate >20/min or PaCO2<32mm Hg
- 3) Body temperature <36'C or >38'C

4) WBC Count <4000 or >12000/mm3 or >10% band forms(immature WBC's)

Sequential Organ Failure Assessment (SOFA) score has been established using systemic failure tests in six organ systems.^[4] (Respiratory, renal, kidney, coagulation and central nervous systems). Everyone is graded as 0 to 4 with an elevated frequency of impairment. Sepsis markers for both C-reactive protein (CRP) and procalcitonin (PCT) are recognized.

Nonetheless, the link between plasma concentrations and septic intensity is still under debate. The plasma amounts of CRP and PCT in sepsis are also measured in a number of severities, as determined by the tests of the Couch.

Subjects and Methods

Type & Place of study: It's a standardized retrospective study was conducted on 50 patients admitted to intensive care units at Gandhi Medical College.

Inclusion Criteria

Patient's \geq 15 years of age admitted to ICU.

Exclusion Criteria

- 1. Patients of age <15 years
- 2. Patients with post-operative and post-traumatic sepsis.

Method of Collection of Data

Patients chosen for the study are grouped into 4 categories depending on the extent of sepsis organ impairment according to the SOFA score.

- Group 1 0-6
- Group 2 7-12
- Group 3 13-18
- Group 4 19-24

Serum PCT and CRP concentrations were estimated in all the patients.

Statistical Methods

Statistical tables and charts were used to illustrate the statistical data. For statistical analysis, SPSS software version 20 was used.

Results

The majority of the patients belonged to the age groups of 60-69 years (30%) and 50-59 years (22%) and the least belonging to the age group of 15 -29, 30-39 years both with (8%) and Age of 80 years and above with 4%.

The majority of the patients belonged to the Sofa group 1 around 42% followed by sofa group 2 with 38%, sofa group 3 with 16% and the least belonged to sofa group 4 with 4%.

Type of Infections seen in sepsis patients with the majority of the patients being diagnosed with Pneumonia around 42% of them followed by Urinary tract infection (UTI) seen in 22% of the patients with sepsis.

The mean PCT was more in SOFA group 4 around 87.71 ng/ml and it was least in group 1 around 7.237 ng/ml. The mean CRP

Table 1: Age & Sex Distribution					
Age Group (Yrs.)	Total No. of Patients (N=50)	Percentage %			
15-29	4	8%			
30-39	4	8%			
40-49	9	18%			
50-59	11	22%			
60-69	15	30%			
70-79	5	10%			
80 & above	2	4%			
Sex					
Male	29	58%			
Female	21	42%			

Table 2: SOFA Groups and Number of Patients					
Total Sofa Score	SOFA Group	Number of patients (n=50)	Percentage %		
0-6	1	21	42%		
9-12	2	19	38%		
13-18	3	8	16%		
19-24	4	2	4%		

Table 3	3: Infections	are se	en in	Patients	Belonging	to	Different
SOFA	Groups						

Infection	Number of patients (n=50)	Percentage %
Pneumonia	21	42%
Urinary tract infection (UTI)	11	22%
Diarrhea	6	12%
Pyogenic menin- gitis	4	8%
Cellulitis	2	4%
Ludwig's angina	1	2%
SBP	1	2%
Dengue	1	2%
Necrotizing fasci- itis	1	2%
Viral meningitis	1	2%
Malaria	1	2%

Table 4: Organ Dysfunction (Sofa Group) Vs Mean Serum PCT and CRP Concentration						
SOFA Score	SOFA Group	Total (N=50)	Percentage %	Mean (ng/ml)	РСТ	Mean CRP (mg/L)
0-6	1	21	42%	7.237		159.53
7-12	2	19	38%	28.172		155.23
13-18	3	8	16%	43.23		146.22
19-24	4	2	4%	87.71		41.73

 Table 5: Outcome Vs Mean Serum PCT and CRP Concentration

Outcome	No. of patients	Percentage %	Mean PCT (ng/ml)	Mean CRP (mg/L)
Survival	41	82%	14.73	149.916
Death	9	18%	45.76	183.584
Total	50	100%	21.89	155.332

was least found in group 4 with 41.73mg/L and highest in the group 159.53 mg/L.

The survival rate of patients was 82% with mean PCT being 14.73 ng/ml and mean CRP being 149.916mg/L in the survived patients. In patients who died around 18%, the mean PCT was found to be 45.76 ng/ml and mean CRP was found to be 183.584mg/L.

With respect to outcome mean SOFA score and serum PCT (ng/ml) were higher with death than with survival (p-value <0.001).

Discussion

The SOFA group mortality rate increased from group 1 to 4 indicating a higher mortality rate with serious and multiple organ dysfunction. SOFA and mean serum PCT concentrations were higher for SOFA (groups 3 and 4) than for patients with mild organ impairment (groups 1 and 2 of SOFA) in our sample (p<0.001). There was no statistically relevant difference in the concentration of mean serum CRP in the SOFA (organ dysfunction) classes. The mean SOFA score (organ dysfunction) was higher in patients who died than in those who had survived. (p < 0.001). The mean PCT (ng/ml) serum concentration was higher for those who died than for surviving patients and the difference was statistically significant (p < 0.001). Mean serum CRP (mg / L) concentration was not statistically relevant to the outcome. The positive association between the SOFA value and the mean PCT serum concentration was statistically important. There were no significant associations in our sample between the SOFA value and the mean PCT concentration. The findings of our analysis indicate that PCT rates were favorable for the extent of sepsis organ impairment as assessed by SOFA. The findings of our study are broadly compatible with studies that compared PCT levels with sepsis frequency with sepsis-

related scoring systems. Previous tests demonstrated elevated rates of PCT and relatively small rates of SIRS or less serious systemic inflammation during a septic shock. Increased PCT concentrations were previously recorded during more extreme septic phases (serious sepsis / septic shock) as described in the ACCP / SCCM guidelines of Oberhofer et al,^[5] and Zeni et al.^[6] Al-Nawas et al,^[7] registered very low PCT concentrations during SIRS, but high concentrations were found when septic shocks were detected. Gramm et.al, [8] and de Werra et.al,^[9] showed identical findings. Such research does not evaluate the extent and intensity of sepsis and chronic inflammation with multiple organ failure. In a retrospective empirical analysis by Pugnani C et.al Castelli GP, PCT and SOFA were more likely to be higher in septic shock than extreme sepsis and sepsis.^[10] They found that PCT and SOFA were most directly linked to the extent of the infection. The results of our study are compatible with this statement. PCT has several advantages over CRP in severely ill sepsis patients. Concentrations of PCT are very weak in moderate organ impairment or bad systemic inflammatory reactions. Nevertheless, CRP levels are still strong for patients with higher SOFA concentrations. As a consequence, relative to PCT, CRP does not have useful statistics on the extent of organ dysfunction because the average rates of organ dysfunction have already been elevated during the more severe stage of the disease. Patients with low prognosis, i.e. patients with extreme organ failure (strong SOFA score) and those with sepsis have recorded elevated PCT rates. The key drawback of our research is the limited sample range in the classification of patients with extreme organ failure (SOFA classes 3 and 4). In comparison, organ impairment was not taken into consideration in the study of the association between serum PCT and CRP concentrations. In order to derive from the absolute amount of PCT concentrations to estimate the PCT prognosis, the number of patients in our study is very low to heterogeneous.

Table 6: Comparison of study variables according to the outcome					
Variables	Survival	Death	P-value		
SOFA Score	6.12±3.40	11.03±4.51	<0.001**		
S.PCT (ng/ml)	14.73±19.51	45.76±37.01	<0.001**		
S.CRP (mg/L)	149.91±56.41	$183.584{\pm}65.05$	0.172		

Conclusion

Patients with extreme organ dysfunction (SOFA group 3&4) have a mean SOFA and PCT value greater than people with moderate organ dysfunction (SOFA group 1&2) (p<0.001). The increase in the SOFA score was associated with a higher median value of PCT, but not of CRP. In patients who died of sepsis, the mean concentration of SOFA Score & PCT was greater than that of survivors. Mean CRP was not substantially associated with the extent of the organ dysfunction/result. The ability of PCT to cause very significant volumes of serum during advanced phases of CRP-based modifications is characterized by its intensity while CRP is still already in its upper concentration range with small rates of SOFA.PCT and SOFA are strongly linked to the extent of the infection. PCT is easier than CRP to assess the frequency and prognosis of sepsis. Owing to the strong association between the PCT stage and the sepsis intensity and effect, the PCT diagnostic performance is superior to the CRP.

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