



Can Biomarkers Help Identifying The Type Of Blood Stream Infection In Septic Patients?

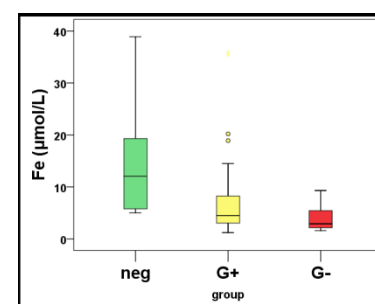
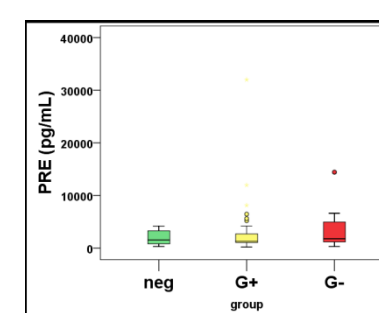
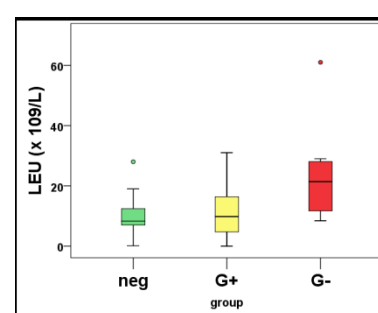
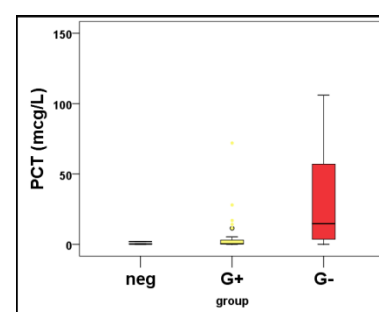
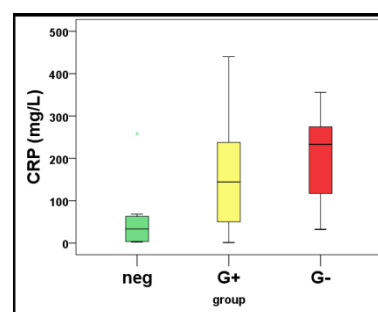
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Introduction: Sepsis is one of the most prevalent causes of morbidity and mortality in hospitalized patients worldwide. Early initiation of targeted antibiotic therapy is crucial [1]. Blood stream infections are commonly divided to Gram positive (G+) or Gram negative (G-). However, blood cultures (BC) read-out may be delayed or cultures may be negative (NEG). This could affect the patient outcome [2,3]. Biomarkers may help to guide antibiotic therapy prior to BC results [4-6]. We tested the hypotheses that 1) biomarkers will discriminate between BC- vs. BC+ patients; 2) biomarkers will discriminate between G+ and G- sepsis; 3) biomarkers will correlate with severity of illness.

Methods: With IRB approval, a patient cohort (n=60) admitted to mixed ICU for suspected sepsis were enrolled in a prospective observational study. BC and biomarkers of sepsis (C-reactive protein, CRP; procalcitonin, PCT; presepsin, PRE; leukocytes, LEU and iron, Fe) were assessed and SOFA and qSOFA were determined on admission. Data are displayed as mean±SD or median [IQR]. One-way ANOVA with post-hoc Tukey's test, Kruskal-Wallis test or Mann-Whitney test were used as appropriate. Pearson's test was used to assess correlation between SOFA and biomarkers.

Results: G- sepsis was more common in older patients and severity of their status was higher than G+ and BC - patients. CRP was the only biomarker different between BC- (33 [3, 64] mg/L) vs. BC+ (147 [51, 256] mg/L) patients (p=0.003). Numerically higher values were observed in G- patients. CRP was higher in G+ (p=0.023) and G- (p=0.006) vs. NEG. PCT was higher in G- vs. NEG (p=0.037). LEU were higher in G- vs. NEG (p=0.044) and vs. G+ (p=0.015). PRE was not different between groups. Fe was lower in G+ vs NEG (p=0.021), in G- vs NEG (p=0.004) with no difference between G+ and G- (p=0.537). In NEG patients, SOFA score did not correlate with any biomarkers. In BC+ patients, SOFA correlated with CRP (p=0.005) and LEU (p=0.023). PRE correlated with SOFA in G+ patients (p=0.032).



	NEG (n=10)	G+ (n=39)	G- (n=11)
age (yrs)	53±18	64±16	77±8†‡
SOFA (pts)	3.9±2.2	4.7±2.2	8.1±4.5
qSOFA (pts)	0.2±0.4	0.6±0.8	0.9±0.9

† p<0.05 vs. NEG; ‡ p<0.05 vs. G+

Conclusion:

In our limited sample-size pilot study, tested biomarkers showed limited capacity to identify BC+ patients and the type of infection. Higher values of inflammatory biomarkers observed in G- sepsis and lower values of Fe in positive BC warrant further study in a larger population of patients.

References:

- [1] Seymour et al. N Engl J Med 2017.
- [2] Kethiredy et al. Crit Care Med 2018 (epub)
- [3] Gupta et al. Chest 2016.
- [4] Brodska et al. Clin Exp Med 2013.
- [5] Leli et al. Disease Markers 2015.
- [6] Weis et al. Cell 2017.