

THE EMERGENCY DIAGNOSTIC MARKERS IN PEDIATRIC PATIENTS WITH SUSPECTED SEPSIS/BACTEREMIA

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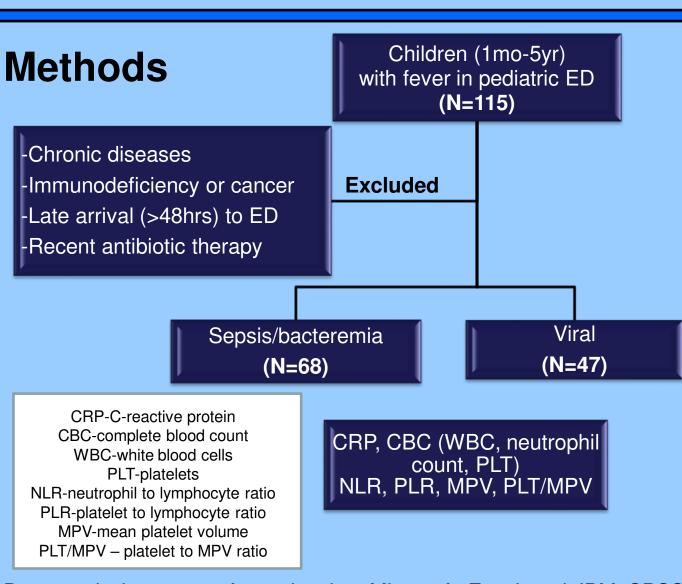


Background

- -Sepsis is a clinical syndrome that is triggered by bacterial infection, which may lead to multiple organ dysfunction (1, 2)
- -Quick and early detection is necessary in order to prevent more severe forms and lethal outcomes
- -Sepsis specific early biomarkers are missing in clinical practice (3)

Aims

- -To investigate the diagnostic value of NLR, PLR, MPV, PLT and PLT/MPV as early markers (<12h after onset of symptoms) in viral infection vs bacterial (sepsis/bacteremia) in pediatric emergency department (ED) settings
- -To apply cut-off levels of inflammatory markers
- -Develop a prediction model to distinguish between severe bacterial and viral infection in pediatric patients



Data analysis was performed using Microsoft Excel and IBM SPSS Statistics version 21.0 software. P value <0.05 was considered significant

Results

	Sepsis/bacteremia	Viral infection	P value
Age (months)	9 [3–24]	12 [6–27]	0.274
Gender (male)	31 (45.6%)	23 (48.9%)	0.857

Table 1. Baseline characteristics for all children, at any time of arrival

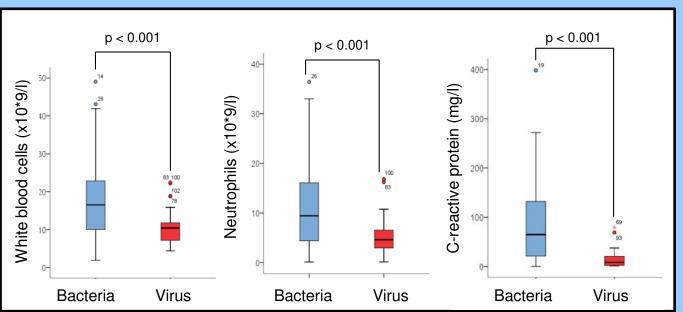


Figure 1. Sepsis/bacteremia vs. viral infection: Comparison of the entire study population, n = 115

- 1.Weiss SL, et al. Global epidemiology of pediatric severe sepsis: the sepsis prevalence, outcomes, and therapies study. Am J Respir Crit Care Med. 2015;191(10):1147-57
- 2.Reinhart, K., et al. New Approaches to Sepsis: Molecular Diagnostics and Biomarkers. Clin Microbiol Rev. 2012; 25(4): 609–634
- 3.Lanziotti, V. S., et al. Use of biomarkers in pediatric sepsis: literature review. *Rev Bras Ter Intensiva. 2016; 28*(4), 472–482

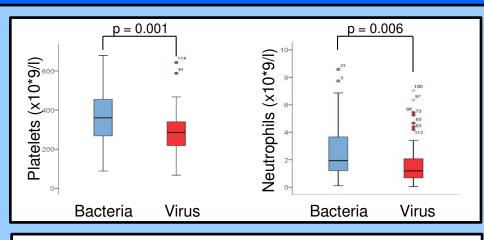


Figure 2.
Comparison of laboratory markers between bacterial and viral infection patients (early and late arrival)

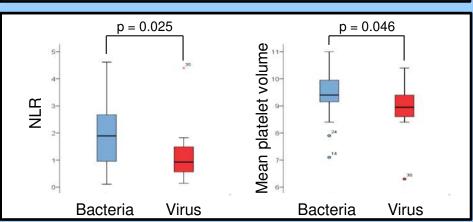


Figure 3.
Sepsis/bacteremia
vs. viral infection:
Infants (≤12 months)
with late (>12 hours)
arrival to ED,
n = 38

Laboratory marker	AU	С	95% C	I	P value
WBC (>11.5 ×10 ⁹ /L)	0.73	30	0.63–0.8	33	0.049
Neutrophils (>5.35 ×10 ⁹ /L)	0.676		0.57–0.78		0.002
PLT (>315 ×10 ⁹ /L)	0.631		0.53–0.74		0.018
CRP (>20 mg/L)	0.748		0.65–0.84		<0.001
NLR (1.58)	0.653		0.55–0.76		0.006
Laboratory marker			ensitivity	Зp	ecificity

PLI (>315 ×10 ⁸ /L)	0.631		0.53-0.74		0.018
CRP (>20 mg/L) 0.74		48	0.65–0.84		<0.001
NLR (1.58)	0.6	53	0.55–0.76		0.006
Laboratory mark (cut-off)	er	Se	ensitivity (%)	Sp	ecificity (%)
WBC (>11.5 ×10 ⁹ /L)		80.3		64.8	
Neutrophils (>5.35 ×10 ⁹ /L)			74.2		61.2
PLT (>315 ×10 ⁹ /L)			71.7		54.5
CRP (>20 mg/L)			80.3		68.8
NLR (1.58)			73.0		57.7

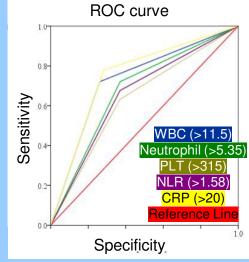
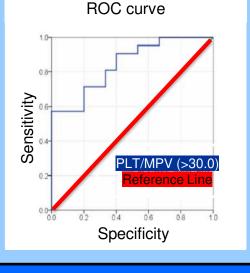


Table 2. Sensitivity and specificity, area under the curve (AUC) and confidence intervals for cut-off values for all children who arrived early to ED (<12 hours)

Laboratory marker (cut-off)	AUC	95% CI	P value
	0.844	0.72–0.97	<0.001
PLT/MPV (30.0)	Sensitivity 95.2	•	ficity (%) 53.3

Table 3. Sensitivity and specificity, area under the curve (AUC) and confidence intervals for cut-off values for **all children** who arrived **early** to ED (<12 hours)



Conclusion

- -WBC and CRP showed to be the **most reliable** biomarkers to determine bacterial infection and differentiate from viral infections, at **any time** of arrival
- -PLT, NLR and PLT/MPV could be considered as a bacterial infection marker in children who arrived to ED early
- -PLT/MPV is one of the most sensitive early bacteremia/sepsis markers in infants