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20-30 JUNE 2021 | **Q** ONLINE



# Correlation of Inflammation, Lipidogram and Clinical Readings in Chronic Heart Failure Patients

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#### Introduction (I)

- Traditional model of heart failure (HF) pathophysiology: HF with reduced ejection fraction (HFrEF) has been mainly attributed to ischemic left ventricular remodeling [1-3], whereas HF with preserved ejection fraction (HFpEF) has been attributed to hypertension [4-7].
- **Novel paradigm** of chronic HF (CHF) pathogenesis: metabolism-related concomitant diseases play a crucial role in systemic pro-inflammatory condition maintenance in HFpEF [5,8-14].



#### Introduction (II)

- Inflammatory processes are presented as regulated by plateletinduced activation of blood leukocytes.
- Neutrophils take part in maintaining a pro-inflammatory state in the pathophysiology of HF [15].
- Hypercholesterolemia is stated to heighten neutrophil production, which contributes to accelerated cardiovascular inflammation [16].
- Identification of inexpensive, reliable, and most importantly, rapid prognostic markers of HF.
- HF pathogenesis differences in different HF phenotypes remain to be investigated.



#### Aim

To determine differences in complete blood count, C-reactive protein (CRP) concentration, lipidogram and clinical readings between CHF without previous MI groups according to EF and between HFrEF groups according to MI presence in CHF development history and correlations between these readings.



#### Methods

- 4 groups of patients were analyzed (n = 266).
- **Period:** from January 1, 2018 to February 1, 2021 (Hospital of Lithuanian University of Health Sciences Kauno klinikos Cardiology department).
- 208 patients diagnosed with CHF who had had no documented history of previous myocardial infarction (MI) were divided into two groups according to left ventricular ejection fraction (LVEF): LVEF ≥ 50%, n = 117 and LVEF < 50%, n = 91.
- Additionally, 149 HFrEF patients were separated into two additional groups: those who had had no MI (n = 91) and those with MI (n = 58).
- Exclusion criteria: malignancies, chronic obstructive pulmonary disease, bronchial asthma, autoimmune diseases, stage 4–5 chronic kidney disease (CKD, with eGFR < 30 ml/min/1.73²), acute infections, i.e., common chronic or acute systemic inflammation supporting conditions.

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## Results (I)

Laboratory findings	LVEF ≥ 50 % without MI, n=117	LVEF < 50 % without MI, n=91	p - value
RBC, 10 <sup>12</sup> /l	4.59 (0.57)	4.61 (0.65)	0.791
HGB, g/I	137 (87-165)	136 (77-183)	0.477
MCHC, g/I	337.32 (10.60)	331.46 (13.13)	0.004*
PLT, 10 <sup>9</sup> /l	202 (73-326)	204.5 (113-1097)	0.053
RDW-CV, %	13.6 (11.5-16.9)	14.7 (12.6-19.1)	0.001*
MPV, fl	9.85 (1.37)	10.13 (1.19)	0.222

LVEF – left ventricular ejection fraction, MI – myocardial infarction, RBC – red blood cells, HGB – hemoglobin concentration, MCHC – mean corpuscular hemoglobin concentration, PLT – platelets, RDW-CV – red cell distribution width, MPV – mean platelet volume \* Statistically significant values (p < 0.05)



### Results (II)

Laboratory findings	LVEF ≥ 50 % without MI, n=117	LVEF < 50 % without MI, n=91	p - value
NEU, %	58.20 (12.40)	61.12 (10.40)	0.137
NEU, 10 <sup>9</sup> /I	4.00 (1.42-15.53)	4.05 (1.47-9.61)	0.434
LYM, %	30.48 (10.87)	26.98 (9.08)	0.045*
LYM, 10 <sup>9</sup> /l	1.98 (0.72)	1.78 (0.59)	0.071
MON, %	9.1 (4.7-13.7)	9.4 (3.2-15.9)	0.101
MON, 10 <sup>9</sup> /I	8.78 (2.69)	9.52 (2.81)	0.121
LYM/MON	3.33 (1.22-9.33)	3 (0.44-6.5)	0.011*
CRP, mg/l	4.92(6.21)	7.51(12.29)	0.099

LVEF – left ventricular ejection fraction, MI – myocardial infarction, NEU – neutrophils, LYM – lymphocytes, MON – monocytes, LYM/MON – lymphocyte-to-monocyte ratio, CRP – C-reactive protein concentration



<sup>\*</sup> Statistically significant values (p < 0.05)

## Results (III)

Laboratory findings	LVEF < 50 % without MI, n=91	LVEF < 50 % with MI, n=58	p - value
Total cholesterol, g/l	4.35 (2.46-7.10)	3.9 (2.72-6.71)	0.016*
LDL, g/l	2.97 (1.53-5.5)	2.52 (1.36-4.42)	0.101
HDL, g/l	0.96 (0.44-2.2)	0.92 (0.56-1.97)	0.010*
TG, g/I	1.25 (0.39-3.28)	1.24 (0.51-6.78)	0.672
AC	3.55 (1.23-6.06)	3.25 (1.21-6.39)	0.591
CRP, mg/l	6.9 (1.46-62.97)	7 (1-33.99)	0.012*

LVEF – left ventricular ejection fraction, MI – myocardial infarction, LDL – low-density lipoprotein concentration, HDL – high-density lipoprotein concentration, TG – triglyceride concentration, AC – atherogenic coefficient, CRP – C-reactive protein concentration \* Statistically significant values (p < 0.05)



#### Results (IV)

- Neutrophil count correlated with PLT (rs=0.278, p=0.001) and weight (rp=0.196, p=0.024).
- Lymphocyte count correlated with PLT, RDW-CV ( $r_s$ =0.200, p=0.018;  $r_s$ =-0.223, p=0.032) and body mass index ( $r_p$ =0.186, p=0.032).
- RDW-CV and monocyte count correlated with NT-proBNP and serum creatinine (rs=0.358, p=0.034; rs=0.424, p<0.001 and rs=0.354, p=0.012; rs=0.205, p=0.018 respectively).



#### Results (V)

- Total cholesterol concentration correlated with lymphocyte-to-monocyte ratio (LYM/MON), monocyte percentage, lymphocyte percentage and count (rs=0.534, p<0.001; rs=-0.312, p=0.029; rs=0.355, p=0.012; rs=0.397, p=0.004 respectively).
- LVEF correlated with MCHC and RDW-CV (rs=0.273, p=0.001; rs=-0.404, p<0.001).



#### Conclusion

- 1. MCHC and lymphocyte percentage were lower and RDW-CV was higher in the HFrEF group without of MI; CRP concentration was higher in HFrEF with MI in comparison with the group without MI;
- 2. HDL cholesterol concentration was lower and CRP concentration was higher in the HFrEF group with MI in comparison with the group without MI; total cholesterol concentration correlated with LYM/MON;
- 3. Monocyte, lymphocyte count and their ratio correlated with patients' condition reflected readings NT-proBNP, serum creatinine, uric acid concentrations.

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