

## NEUTROPHIL-TO-LYMPHOCYTE RATIO AND MIR-126-3P AS BIOMARKERS OF NOCTURNAL AND RESISTANT HYPERTENSION IN CKD PATIENTS.

D Klimczak-Tomaniak<sup>1,2</sup>, T Pilecki<sup>1</sup>, D Żochowska<sup>1</sup>, D Sieńko<sup>1</sup>, M Janiszewski<sup>2</sup>, M Kuch<sup>3</sup>, L Paćzek<sup>1</sup>

1. Department of Immunology, Transplantation and Internal Medicine, Transplantation Institute, Medical University of Warsaw, Warsaw, Poland 2. Department of Heart Failure and Cardiac Rehabilitation, Medical University of Warsaw, Warsaw, Poland. 3. Chair and Department of Cardiology, Hypertension and Internal Medicine, Medical University of Warsaw, Warsaw, Poland.

**Introduction :** Pro-inflammatory milieu of chronic kidney disease (CKD) increases cardiovascular risk of this patient population. Neutrophil-to-lymphocyte ratio (NLR) is an easily available inflammatory parameter whereas microRNA-126-3p is regarded as a novel biomarker of endothelial damage.

**Objectives:** The aim of the study is to evaluate the association between NLR and relative expression of miR-126-3p with blood pressure values and circadian blood pressure (BP) abnormalities in CKD patients.

**Patients and methods:** This single-centre observational study involved stable CKD stage 1-5 patients and healthy age- and sex- matched control subjects with no clinical evidence of infection. All study participants had 24-hour automatic blood pressure measurement (ABPM) performed. Plasma miRNA was quantified by qRT-PCR with the use of the  $\Delta\Delta C_t$  method and endogenous control U6 snRNA.

**Results:** In total, 90 CKD patients ( $60 \pm 14$  years, 52% males, 33 renal transplant recipients) and 25 healthy control subjects ( $55 \pm 13$  years, 48% males,  $p < 0.05$ ) were enrolled in the study. Subjects with nocturnal hypertension (N=41, 46%) presented with higher plasma relative expression of miR-126-3p ( $4.0 [2.2-7.0]$  vs.  $1.7 [1.0 - 3.5]$ ,  $P = 0.004$ ) compared to patients with normal night-time BP values. We have observed a positive correlation between plasma miR-126-3p and average nocturnal SBP ( $\rho = 0.27$ ,  $P = 0.02$ ), average nocturnal DBP ( $\rho = 0.32$ ,  $P = 0.003$ ), night-day SBP ratio ( $\rho = 0.23$ ,  $P = 0.03$ ), and night-day DBP ratio ( $\rho = 0.26$ ,  $P = 0.02$ ). A positive association was found between NLR and average nocturnal SBP ( $\rho = 0.25$ ,  $P = 0.01$ ), ND-SBP ratio ( $\rho = 0.26$ ,  $P = 0.006$ ) and ND-DBP ratio ( $\rho = 0.28$ ,  $P = 0.03$ ). Based on the multivariate regression analysis that included bivariate correlates of nocturnal SBP ( $P = 0.1$ ), NLR was the the only independent predictor of nocturnal SBP ( $B = 11.2$ , 95%CI 1.8 – 10.6,  $P = 0.02$ ). NLR was higher in patients with resistant hypertension (N=20, 22%), being the only biochemical parameter that distinguished this group from the rest of the study cohort ( $3.0 [2.3 - 4.2]$  vs.  $2.0 [1.6 - 2.9]$ ,  $P = 0.01$ ).

**Conclusions:** Increased NLR may reflect the involvement of inflammatory pathways in the development of nocturnal and resistant hypertension. MiR-126-3p may serve as a biomarker of endothelial damage in CKD population with abnormal circadian blood pressure pattern.