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### **P1.27: NT-PROBNP AND VASCULAR CALCIFICATION IN AFRICAN AND CAUCASIAN MEN: THE SAFREIC STUDY**

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unadjusted results analysed by unpaired t-test, shows that obese individuals have a higher aPWV of 0.2007 m/s compared with the control group ( $P = 0.0338$ , 95% CI: 0.39 to 0.02). When analysing data in correspondence with a European normal material obese children had an aPWV standard deviation score in the normal range for age and height although higher than the control group in the present study.

This study shows that obesity is correlated to a higher aPWV although this effect may disappear after adjustment for possible confounders.

#### P1.27

##### NT-PROBNP AND VASCULAR CALCIFICATION IN AFRICAN AND CAUCASIAN MEN: THE SAFREIC STUDY

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**Background:** The N-terminal prohormone B-type natriuretic peptide (NT-proBNP) is a reliable marker of cardiac strain. In hypertensive heart disease, NT-proBNP levels increase and may lose its protective function. Simultaneously, the vasculature is also subject to hemodynamic stress, resulting in vascular matrix remodelling and stiffening which contribute to further cardiac alterations. Alkaline phosphatase (ALP) is a marker of osteoblast activity and is involved in vascular calcification. We explored the link between NT-proBNP and ALP in African and Caucasian men.

**Design and measurements:** This study included 128 African (mean age, 41.1 years) and 118 Caucasian (mean age, 36.4 years) men. Conventional measurements were acquired along with serum NT-proBNP and ALP.

**Results:** NT-proBNP correlated positively with ALP ( $r = 0.29$ ;  $p < 0.001$ ) in Africans, but inversely in Caucasians ( $r = -0.20$ ;  $p = 0.024$ ). After minimal adjustment (age, body mass index, SBP and arterial compliance), the positive significant correlation of NT-proBNP with ALP remained in African men ( $r = 0.225$ ;  $p = 0.014$ ), whereas significance was lost in Caucasian men. Multiple regression analyses confirmed the independent association of NT-proBNP with ALP in African men ( $R^2 = 0.37$ ;  $\beta = 0.248$ ;  $p = 0.005$ ), as well as in younger African men ( $R^2 = 0.26$ ;  $\beta = 0.375$ ;  $p < 0.001$ ;  $n = 96$ ), with no significance in Caucasians.

**Conclusions:** NT-proBNP is independently and positively associated with ALP in African men. This was however not evident in Caucasian men. These results suggest that African men are susceptible to early vascular calcification and may develop cardiac afterload prematurely.

#### P1.28

##### THE ACCUMULATION OF RISK FACTORS OF METABOLIC SYNDROME IS ASSOCIATED WITH THE INCREASE IN ARTERIAL STIFFNESS AMONG MIDDLE-AGED MALE INDUSTRIAL WORKERS

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Early detection of arterial dysfunction is important in preventing cardiovascular diseases. We evaluated the associations of the accumulated cardiovascular risk factors with the cardio-ankle vascular index (CAVI), a new indicator of arterial stiffness.

**Methods:** The study population consisted of 101 employees in the forest industry (49±8 years of age). CAVI was measured by PWV method (VaSera, Fukuda Denshii, Japan). International Diabetes Association (IDF) criteria for metabolic syndrome were used to dichotomize the cardiovascular (CV) risk variables (<http://www.idf.org/metabolic-syndrome>). Statistical analyses were performed by SPSS 20.0 for Windows (SPSS, USA). Number of risk factors was used as a grouping variable for group comparisons. Independent-Samples T Test was used to compare means and stepwise multiple regression to evaluate the independent risk factors affecting CAVI.

**Results:** There were no differences between number of risk factors and CAVI in women. Only the men ( $N = 72$ , 49±8 years of age) were included in further analyses. The CAVI of the men with 3 or 4 risk factors was significantly higher than the CAVI of the men without risk factors ( $p$ -values 0.025 – 0.0005). Men with elevated blood glucose level (B-Gluc) or arterial blood pressure (BP) had significantly higher CAVI than men with normal B-Gluc ( $p = 0.032$ ) and BP ( $p = 0.007$ ). In the regression analysis Age ( $\beta = 0.455$ ,  $p = 0.000$ ), B-Gluc ( $\beta = 0.237$ ,  $p = 0.038$ ), systolic blood pressure ( $\beta = 0.268$ ,  $p = 0.010$ ) and

waist circumference ( $\beta = -0.201$ ,  $p = 0.064$ ) explained 45.1 % of the variation in CAVI.

**Discussion:** Among middle-aged men the number of CV risk factors is an important determinant of cardiovascular health assessed by arterial stiffness. B-Gluc and BP may have a special negative effect on CAVI. Waist circumference seems to be more useful risk factor for arterial dysfunction in male workers than BMI.

#### P1.29

##### AN IGF2 RECEPTOR GENE POLYMORPHISM MODULATES BLOOD PRESSURE TRENDS OVER TIME IN TYPE 2 DIABETES

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**Rationale:** Previous studies have associated low circulating IGFBP-2 with diastolic BP in a cross-sectional population. We examined 1) the relation between IGFBP-2 and long term trends in blood pressure in a population with type 2DM and 2) the relation between blood pressure and SNPs from the IGFBP-2 and IGF2 receptor (*IGF2R*) gene.

583 individuals with T2DM (58.5% male;  $n = 341$ ) had repeated yearly cardiometabolic assessments between 2002 and 2009. We used a commercial ELISA (RayBio Inc) platform for IGFBP-2 measurement. Haplotype tagging SNPs (8 from *IGF2* gene, 12 from *IGF2R* gene and 2 from *IGFBP2* gene) were selected.

**Results:** High baseline IGFBP-2 ( $\beta = -1.52$ , 95% CI: -2.56, -0.49,  $p = 0.004$ ) was associated with a longitudinal decrease in diastolic BP over 8 years, adjusted for age, gender, diabetes duration, time effects, as well as IGF-I, IGF-II, IGFBP-1, IGFBP-3 and anti-hypertensive use. There was no association in a similar model using systolic BP. In mixed-effects regression models the SNP rs2014620 from the *IGF2R* gene (encoding the IGF-II receptor which degrades IGF-II) was associated with decreased diastolic BP over the 8-year period adjusted for age and gender ( $\beta = -0.252$ , 95% CI -0.14 to -0.298,  $p = 0.003$ ). Significance remained after gene-wise Bonferroni adjustment. This SNP rs2014620 was also nominally associated with higher baseline IGFBP-2 adjusted for age and gender ( $\beta = 0.119$ ,  $p = 0.011$ ).

**Conclusion:** We suggest that SNPs in the *IGF2R* gene may influence IGF-II bioavailability independently of IGF-II degradation, with the possibility that variations in this gene directly modulate longitudinal diastolic blood pressure trends.

#### P1.30

##### IN HIGH CVD RISK GROUP ARTERIAL FUNCTION MARKERS CORRELATE WEAKLY WITH PWV AND AIXHR75 AS EXCEPTIONS

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**Background:** With more recent publications it is becoming obvious that arterial wall function contributes to the cardiovascular disease (CVD) risk. Relatively few studies investigated high CVD risk subjects, where arterial wall function abnormalities are likely to be more pronounced with multiple risk factors. The aim of the study was to assess the relationship between arterial wall markers in a high CVD risk subjects.

**Methods:** A cross-sectional study included patients with metabolic syndrome but with no previous CVD. Arterial stiffness (aortic pulse wave velocity [PWV] augmentation index [Aix], carotid and ankle-brachial stiffness index [CSI and CAVI, respectively]), endothelial function (flow-mediated dilatation in brachial artery and finger [RHI]), and carotid intima-media thickness (CIMT) were measured. Univariate and multivariable association assessment between these parameters was performed.

**Results:** Among 3168 subjects (aged 55±5, 69% women) univariate analysis revealed that markers significantly ( $p < 0.01$ , Pearson  $r > 0.1$ ) associated with PWV and Aix are CAVI and RHI, and PWV correlates with Aix. CSI was significantly associated only with CIMT. No significant interaction between other independent variables was observed. Interestingly, 2/3 of markers revealed higher correlations for male compared to female group. Traditional risk factors (gender, age, blood pressure, BMI etc.) explained only 12-34% of variability for PWV, Aix, and CSI.

**Conclusion:** In this high CVD risk group we did not find a definite/strong correlation between most of the arterial markers investigated, possibly because they reflect different stage of the same process or due to varying impact of different factors. Stronger inter-correlation of the arterial markers was observed in men.