P9.5: VASCULAR STRUCTURE AND FUNCTION AND HEMODYNAMICS ARE NOT ALTERED IN NORMAL-TENSION GLAUCOMA AT REST


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P9.4 COMPARISON OF DIFFERENT METHODS OF VASCULAR PHENOTYPING IN PATIENTS AT CARDIOVASCULAR RISK

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Objective: Discriminating between patients at high and low cardiovascular risk can be difficult. Screening for the presence of subclinical organ damage may provide additional benefits in predicting cardiovascular events. We aimed to assess the agreement of markers of organ damage with traditional risk scoring methods.

Methods: We performed a comprehensive analysis of vascular health on a cohort of 50 patients recruited from Blood Pressure and Cardiovascular Risk Factor clinics. This included pulse wave analysis, pulse wave velocity (PWV), carotid intima-media thickness (cIMT), brachial flow-mediated dilation and ankle-brachial pressure index. Patients were subdivided according to the ASSIGN score into low, intermediate and high cardiovascular risk.

Results: PWV (8.63, 8.25 and 11.9 m/s) and cIMT (0.67, 0.73 and 0.91 mm) were the only vascular parameters to display a statistically significant (p < 0.001) difference according to risk category (low, intermediate and high). Both correlated with age (r = 0.589, p < 0.001; and r = 0.646, p < 0.001; respectively) and with each other (r = 0.585, p < 0.001). When we calculated means and 95% confidence intervals for each risk category we identified 6 low risk, 11 intermediate risk and 6 high risk patients who had test values outwith the ranges expected of their group. For patients with higher or lower values than expected according to their risk category, which could reflect higher or lower risk than originally estimated. These findings require prospective studies.

P9.5 VASCULAR STRUCTURE AND FUNCTION AND HEMODYNAMICS ARE NOT ALTERED IN NORMAL-TENSION GLAUCOMA AT REST

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Background: Increased blood pressure variability (BPV) is associated with adverse cardiovascular outcomes, particularly incident stroke. However, the impact of increased BPV on target-organ damage related to hypertension is unknown. This study aimed to determine the effect of BPV compared with conventional average BP on left ventricular mass index (LVMi).

Methods: BPV and conventional average BP (24-hour ambulatory BP, 7-day home BP and office BP) were assessed in 286 patients with uncomplicated treated hypertension (aged 64±8 years, 54% male) over 12-months. Short-term BPV (from 24 hour ambulatory BP) was assessed at baseline and 12-months, mid-term BPV (from 7 day home BP) was assessed at 3-month intervals, and long-term BPV (from clinic BP) was assessed from 5 visits over 12-months. Left ventricular mass index (LVMi) was derived from 3-dimensional echocardiography.

Results: Mean changes in LVMi over 12 months were 0.01±2.5 g/m².[7]. The strongest independent predictor of the change in LVMi was conventional 24-hour ambulatory systolic BP (β = 0.032; P = 0.02). However, none of the changes in short-term, mid-term or long-term BPV were associated with changes in LVMi (all P > 0.05).

Conclusion: The change in conventional average 24-hour ambulatory BP, but not BPV, is relevant to changes in target organ damage determined from LVMi in patients with uncomplicated hypertension. Thus, BPV does not provide additive information on BP control in this patient population.

P9.6 BLOOD PRESSURE VARIABILITY AND TARGET ORGAN DAMAGE IN PATIENTS WITH UNCOMPPLICATED HYPERTENSION: AVERAGE 24 HOUR AMBULATORY BLOOD PRESSURE IS MORE RELEVANT TO CHANGES IN LEFT VENTRICULAR MASS INDEX

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Background: Blood pressure (BP) variability (BPV) has been shown to be associated with adverse cardiovascular outcomes. However, the extent to which BPV impacts on target organ damage (TOD) in patients with uncomplicated hypertension is unclear.

Methods: 86 patients (mean age 59±10 years, 41 women) with recently diagnosed (median 1.8 years) type 2 diabetes and 86 controls matched individually for gender and age were included. Radial artery pressure waveforms were obtained non-invasively by applanation tonometry. The central aortic waveform was derived using the Sphygmoman transfer function, which enabled calculation of SEVR. Arterial stiffness was assessed by the carotid-femoral pulse wave velocity (PWV).

Results: SEVR was significantly lower in diabetic women compared with i) diabetic men (161% vs. 178% ± 32%, p < 0.001), ii) non-diabetic women (185% ± 24%) and men (188% ± 28%, p < 0.001 for both comparisons). The differences remained significant in a multivariate analysis including age, mean blood pressure, heart rate, smoking, total cholesterol and PWV.

Conclusion: SEVR was significantly lower in diabetic women as compared with both diabetic men and non-diabetic subjects. This association was not mediated by arterial stiffness. Low SEVR may independently contribute to the increased cardiovascular morbidity seen in diabetic women.

P9.7 C- REACTIVE PROTEIN, CYSTATIN C AND ARTERIAL STIFFNESS IN RENAL TRANSPLANT RECIPIENTS

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Low grade inflammation renal transplanted (Tx) patients enhances atherogenesis. Cystatin C (CysC) is a sensitive marker of glomerular filtration rate, furthermore inhibits cysteine proteases, therefore anti-atherogenic. Previous studies found elevated C-reactive protein (CRP) and CysC levels in renal transplant recipients. Arterial stiffness parameters such as aortic augmentation (index Aix) and pulse wave velocity (PWV) are established markers of cardiovascular mortality in these patients. However, the association between inflammatory markers, CysC levels and arterial stiffness has not been studied. 131 Tx patients and 63 age- and gender-matched healthy controls (C) were enrolled in the study. Lipid parameters, CRP and CysC serum levels were measured. Arterial stiffness parameters (Aix, PWV, pulse pressure (PP), systolic and diastolic area indexes (SAI and DAI) and mean arterial pressure (MAP) were determined by arteriograph (Tensionomed). Significantly higher levels of CRP and CysC levels were found in patients compared to controls. Significantly higher MAP, PP, Aix and PWV, while significantly lower DAI and SAI were detected in Tx patients compared to C subjects. Significant positive correlation was found between CRP and MAP, and between CRP and MAP, while there was a significant negative correlation between CRP and DAI in TX patients. We found a non-significant correlation between CysC and PWV. Significant positive correlation was showed between CRP and LDL-C levels.