



## **Artery Research**

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## P1.13: ETHNIC DIFFERENCES IN WAVE INTENSITY AND ARTERIAL STIFFNESS IN THE CAROTID ARTERY

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154 Abstracts

**Conclusions:** In line with experimental studies in animals and high heritability of carotid IMT, we demonstrated that IMT was associated with genetic variations in several interleukins components.

P1.11
CENTRAL SYSTOLIC AUGMENTATION INDEXES AND URINARY SODIUM IN A WHITE POPULATION

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**Background:** The association between cardiovascular health and salt intake remains controversial.

Methods: In 630 participants (mean age 40.6 years; 51% women), randomly recruited from F. alemish population, we measured sodium and creatinine in 24 hour urine samples at baseline and follow-up (median, 9.7 years) and the carotid and aortic augmentation indexes (AI) standardized to heart rate at follow-up only.

**Results:** The carotid AI (130.2% vs 113.7%) and aortic AI (145.7 vs 127.4) were higher (P<0.0001) in women than men and increased with age (10.1% and 8.5% per 10 years). From baseline to follow-up, the urinary sodium concentration decreased (117.1 vs 105.2 mmol/L; P<0.0001), whereas 24 hour urinary sodium did not change (166.5 vs 171.5 mmol; P=0.12). In multivariable-adjusted longitudinal analyses, a 40 mmol/L ( $\sim$ 1 SD) increase in the urinary sodium concentration at baseline was independently and inversely associated with the carotid AI (effect size,  $1.38\pm0.66\%$ ; P=0.038) and aortic AI (1.54 $\pm0.72\%$ ; P=0.019). In cross-sectional analyses of follow-up data, these estimates were  $1.26\pm0.70\%$  (P=0.07) and  $1.52\pm0.76\%$  (P=0.045), respectively. In the longitudinal and cross-sectional analyses, the carotid and aortic AIs were unrelated to the 24 hour urinary excretion of sodium (P>0.43).

Conclusions: Our study showed an inverse association between the Als in the central arteries and the urinary sodium concentration, but not sodium excretion. Vasodilatation of the afferent renal arterioles in response to higher sodium concentration is mediated via the connecting tubule glomerular feedback mechanism; this might move reflection sites in the renal arteries more distally and thereby explain our observations.

P1.12
ETHNIC DIFFERENCES IN LEFT VENTRICLE MYOCARDIAL OXYGEN DEMAND

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**Background:** There are marked inter-ethnic differences in coronary heart disease (CHD). Indian Asians (IA) have 50% greater and African Caribbeans (AC) 50% less CHD than white Europeans (E) in the UK. Reasons for this are unclear. We compared ventricular structure and function, specifically myocardial oxygen demand, by ethnicity.

Methods and Results: 3D echocardiography (Philips iE33) and radial applanation tonometry (SphygmoCor) were performed on 800 men and women (age 55-85) from the Southall And Brent REvisited (SABRE) tri-ethnic population-based cohort. Left ventricular mass index (LVMI) was measured, and 3D LV remodelling index (LVRI) was calculated as LV mass/LV end diastolic volume. 3D cardiac output (CO) and total peripheral resistance (TPR) were calculated and 3D LV end systolic active fibre stress (AFS) and wasted effort (E<sub>w</sub>) were derived as markers of myocardial oxygen demand.

LVMI did not differ between E and AC but was significantly lower in IA. LVRI was greatest in AC and smallest in IA. IA and AC had lower CO and higher TPR compared to E. AFS and  $E_{\rm w}$  were significantly higher in IA. These ethnic differences persisted after multivariate adjustment for age, sex, heart rate, systolic blood pressure, fasting blood glucose and insulin concentrations and medication.

**Conclusions:** AC have comparable LVMI and myocardial oxygen demand to E. In contrast IA generate significantly more AFS and  $E_{\rm w}$  despite having less myocardial muscle. This implies that IA have increased myocardial oxygen demand which may increase susceptibility to myocardial ischemia, and which could contribute to their excess risk of CHD.

**Table 1** Data are mean $\pm$ SE by ethnicity (adjusted for age). \* = p<0.05 \*\* = p<0.01 compared with Europeans by post hoc test following ANOVA.

|  | European         | Indian Asian    | African<br>Caribbean | ANOVA<br>P value |
|--|------------------|-----------------|----------------------|------------------|
| n  | 372              | 294             | 134                  |                  |
| LVMI (g/m <sup>2.7</sup> )                               | $29.7 \pm 0.3$   | 28.2±0.4*       | $29.6 \pm 0.6$       | 0.02             |
| LVRI   | $1.52 \pm 0.02$  | $1.48 \pm 0.02$ | 1.60±0.03*           | 0.005            |
| CO (L)   | $3.5 {\pm} 0.04$ | 3.1±0.05**      | 3.01±0.07**          | < 0.0001         |
| TPR(mmHg/L)  | $29.5 \pm 0.5$   | 33.4±0.5**      | 36.1±0.8**           | < 0.0001         |
| AFS(kPa)   | $22.6 \pm 0.5$   | 24.3±0.6*       | $22.2 \pm 0.8$       | 0.04             |
| E <sub>w</sub> (dyne/s/cm <sup>2</sup> 10 <sup>2</sup> ) | 45±1             | 52±2*           | 49±3                 | <0.0001          |

## P1.13 ETHNIC DIFFERENCES IN WAVE INTENSITY AND ARTERIAL STIFFNESS IN THE CAROTID ARTERY

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Background: In comparison with Europeans (E) in the UK, Indian Asian (IA) people have a 1.5 to 2-fold elevated risk of cardiovascular disease (CVD), and the risk of stroke is more than 2-fold higher in African Caribbean (AC) people. Arterial stiffness and wave reflections influence the development of CVD. We therefore investigated whether there are ethnic differences in wave intensity and arterial stiffness that might play a role in the increased risk of CVD.

Methods and Results: 260 participants, aged 59-82years in the Southall And Brent REvisited (SABRE) population-based study had wave intensity analysis and measurement of stiffness index ( $\beta$ ) performed in the left common carotid artery using an Aloka SSD 5500 ultrasound system (ALOKA, Japan) equipped with a 7.5MHz linear array vascular probe and a combined colour Doppler and echo-tracking system. The intensity of the forward compression wave (FCW) due to left ventricular ejection was significantly increased in IA. The peak intensity of the reflected (backward) compression wave (BCW) was significantly larger in AC.  $\beta$  was significantly higher in both IA and AC. The ethnic differences in the FCW and  $\beta$  persisted after adjustment for key CVD risk factors (Model 2).

**Conclusion:** Both IA and AC have adverse wave intensity and arterial stiffness patterns, independent of conventional CVD risk factors, that may contribute to the increased risk of CVD in IA and AC.

|                         | European<br>n=112 | Indian Asian<br>n=97 | African<br>Caribbean<br>n=51 | ANOVA P  |  |  |
|-------------------------|-------------------|----------------------|------------------------------|----------|--|--|
| FCW (W/m <sup>2</sup> ) |                   |                      |                              |          |  |  |
| Model 1                 | $9.06{\pm}0.05$   | 9.27±0.05**          | $9.05{\pm}0.07$              | 0.009    |  |  |
| Model 2                 | $9.06{\pm}0.05$   | 9.23±0.05**          | $9.04{\pm}0.07$              | < 0.0001 |  |  |
| BCW (W/m <sup>2</sup> ) |                   |                      |                              |          |  |  |
| Model 1                 | $3.48{\pm}0.05$   | $3.58{\pm}0.06$      | 3.66±0.08*                   | 0.1      |  |  |
| Model 2                 | $3.49{\pm}0.06$   | $3.56{\pm}0.06$      | $3.68{\pm}0.08$              | 0.4      |  |  |
| β                       |                   |                      |                              |          |  |  |
| Model 1                 | $2.21 \pm 0.04$   | 2.34±0.04*           | 2.50±0.06**                  | < 0.0001 |  |  |
| Model 2                 | $2.22{\pm}0.04$   | $2.33{\pm}0.04$      | 2.47±0.06**                  | < 0.0001 |  |  |

Data are presented as mean $\pm$ SE and ANCOVA was performed to examine differences between ethnic groups; \* = p < 0.05, \*\* = p < 0.01. Model 1: adjusted for age. Model 2: adjusted for age, sex, heart rate, height, smoking status, diabetes, hypertension and CVD.