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P157: AORTIC CALCIFICATIONS AND INFLAMMATION ARE ASSOCIATED WITH IN-HOSPITAL COMPLICATIONS IN ACUTE CORONARY SYNDROME

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Objective: Carotid-femoral Pulse Wave velocity (cfPWV), the gold standard for measuring stiffness, is a marker of organ damage (OLD). Even though cfPWV correlates with casual (BPc), central (CBP) and ambulatory (ABPM) blood pressure (BP), evidence is limited for resistant hypertension (RH).

Method: Thirty-three patients (age, 56.1 \pm 8.2 years; weight, 78.0 \pm 12.4 kg; height, 1.62 \pm 0.08 m) with RH participated in a cross-sectional study. Outcomes included clinical data, BPc, ABPM, and carotid-femoral, cfPWV. Correlation analysis was conducted to assess the association between variables; independent t-tests were conducted to compare variables between those participants with cfPWV < and \geq 10 m/s.

Results: Patients (20 women and 13 men) presented a peripheral systolic and diastolic BPc of 144.0 \pm 3.8 mmHg and 82.0 \pm 1.9 mmHg, respectively. The cfPWV correlated with age (r = 0.356, p = 0.045), 24 h systolic BP (24 h SBP) nightime pulse pressure (night PP), 24 h pulse pressure (24hPP), casual systolic (SBPc) and diastolic BP (DBPc), central systolic (CSBP), diastolic (CDBP) and central pulse pressure (CPP); controlled for age the correlation remained significant for 24h SBP (r=0.446, p=0.009) 24hPP (r=0.464, p=0.007), nightPP (r=0.365, p=0.036), SBPc (r=0.620, p<0.001), DBPc (r=0.488, p=0.004), PPc (r=0.592, p<0.001), central SBP (r=0.587, p<0.001), central DBP (r=0.487, p=0.001) and central PP (r=0.506, p=0.003). Patients with lower values of cfPWV (n=26) showed lower SBPc (142.8 \pm 15.9 vs. 162.6 \pm 30.9 mmHg, p=0.025), central SBP (136.0 \pm 15.7 vs. 154.1 \pm 31.8 mmHg, p=0.041) and PP (49.6 \pm 9.5 vs. 60.9 \pm 20.8 mmHg, p=0.043) than patients with cfPWV≥10 m/s (n=7).

Conclusion: Our data shows that cfPWV correlates with SBPc, 24hSBP, 24hPP and CSBP, after controlled for age, in patients with RH.

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DETERMINANTS OF PULSE WAVE VELOCITY IN CHILDREN

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Background: Arterial stiffening measured by Pulse Wave Velocity (PWV) predicts cardiovascular events and mortality in adults. It advances with age and seems accelerated in children with certain disease conditions such as chronic kidney disease or diabetes. The aim of this study was to determine factors that influence PWV in children.

Methods: PWV was captured in 285 children aged 10-14 years attending a Portuguese school. The effects of sex, age, height, weight, body mass index, waist circumference, blood pressure, heart rate and sodium excretion in 24-h urinary samples in PWV were tested.

Results: PWV correlated positively with age, height, systolic blood pressure, diastolic blood pressure and heart rate in males and females (p < 0.05) and with weight in males (p < 0.05). Major predictors for PWV in a multivariate regression analysis were gender, height, weight, diastolic blood pressure, heart rate and body mass index.

Conclusion: Our study found several determinants of PWV in children, some of them modifiable and interfering with cardiovascular outcomes. Future research may provide clarity to the association between PWV in children and cardiovascular events in adulthood.

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CARDIO ANKLE VASCULAR INDEX (CAVI) AS ARTERIAL STIFFNESS MARKER IN SUBJECTS WITH ANKYLOSING SPONDYLITIS

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Background: Ankylosing spondylitis (AS) is a chronic, inflammatory disease of the axial spine that can manifest with various clinical signs and symptoms¹.Cardio-ankle vascular index (CAVI), which is calculated based on the stiffness parameter thus obtained, is theoretically independent of changes in blood pressure. With this distinct advantage, CAVI has been widely applied clinically to assess arterial stiffness in subjects with or without known cardiovascular diseases².

Objectives: The aim of this study was to evaluate the Cardio Ankle Vascular Index (CAVI) in subjects with ankylosing spondylitis pared with controls free of morbidities.

Methods: We enrolled 41 participants in this study. Eighteen patients with diagnosed AS and 23 controls free of comorbidities. CAVI was measured by VaSera VS-1000 (Fukuda- Denshi Company, Ltd, Tokyo, Japan).

Results: The results are expressed as mean \pm standard deviation for continuous variables. The data were analyzed using SPSS v. 24 (SPSS Inc., Chicago, IL). The normality of the data was evaluated with Shapiro-Wilk test. A two-tailed p < 0.05 was considered statistically significant. Individuals with AS exhibited greater pSBP (p < 0.01), DBP (p < 0.05), and MBP (p < 0.01) compared to controls. Moreover, in the AS group we observed a higher CAVI with a mean difference of 1.14 (p < 0.01, 95% CI of .41 to 1.8) (Figure 1).

Conclusion: AS is a chronic inflammatory disease that primarily affects the articular joints of the spine. Individuals with ankylosing spondylitis showed increased CAVI, this contributes to explain the higher risk of cardiovascular disease in this pathological condition.



Figure 1. Cardio-ankle vascular index (CAVI) in patients with ankylosing spondylitis (AS) compared to controls

References

1. Van der Heijde D, Ramiro S, Landewé R et al.: 2016 update of the ASAS-EULAR management recommendations for axial spondyloarthritis. Ann Rheum Dis 2017; 76: 978-91.

2. Shirai K, Utino J, Otsuka K, Takata M. A novel blood pressure-independent arterial wall stiffness parameter; cardio-ankle vascular index (CAVI). J Atheroscler Thromb. 2006;13(2):101–107

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AORTIC CALCIFICATIONS AND INFLAMMATION ARE ASSOCIATED WITH IN-HOSPITAL COMPLICATIONS IN ACUTE CORONARY SYNDROME

Konstantia-Paraskevi Gkini, Dimitrios Terentes-Printzios, Charalambos Vlachopoulos, Iosif Koutagiar, Angeliki Rigatou, **Purpose/Background/Objectives:** Aortic calcifications and inflammation are independent predictors of adverse cardiovascular events. We sought to investigate the association of aortic calcifications and inflammation with in-hospital morbidity and mortality of patients with acute coronary syndrome (ACS).

Methods: Two hundred patients (mean age 66 ± 15 years, 150 males) admitted to our Hospital with ACS from 2016-2017 were included in the study. The extent of aortic arch calcification (AAC) on a postero-anterior plain chest X-ray was divided into four grades (0 to 3). Grades 0 to 1 and grades 2 to 3 were categorized as lower and higher AAC grade respectively. High-sensitivity C-reactive protein (hsCRP) was also assessed. In-hospital complications that included reinfarction, arrhythmias, heart failure, stroke, mechanical complications, renal failure, surgery and death were assessed in all patients.

Results: The majority of patients (n = 132, 66%) presented with non-ST elevation ACS, whereas 68 patients as ST-elevation myocardial infarction (STEMI) (n = 68, 34%). Seventy-seven (38.5%) patients presented with one or more in-hospital complications (6 of them died). Higher AAC grade was visible in 44 patients (22%). Patients with higher AAC had increased risk (Odds ratio [OR] = 2.29, 95% Confidence intervals [CI] 1.03 to 5.12, p = 0.043) for in hospital complications after adjusting for age, gender, STEMI/NSTE-ACS diagnosis (OR = 4.10, 95% CI 2.08 to 8.05 for STEMI diagnosis, p < 0.001) and hsCRP (OR = 1.80, 95% CI 1.10 to 2.93, p = 0.02). **Conclusions:** Our study shows that simple tools can be used to assess the inhospital risk of ACS patients. It also highlights the prognostic role of arterial stiffness and low-grade inflammation in ACS.

P158 ARTERIAL STIFFNESS IN THE VERY OLD: THE AGA@4LIFE RESEARCH PROJECT

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Objective: To study the determinants of Arterial Stiffness (AS) in the elderly. Design and method: Cross-sectional, observational study of elderly participants. Blood Pressure (BP) and arterial function parameters were measured with a validated device. Clinical and demographic information was gathered, as well as the estimation of global cardiovascular risk, health related quality of life, dietary profile and cognition. Cholesterol and glycaemia were measured. Results: 54 Participants recruited for the project, with a mean age of 73.0 \pm 6.0 years (range: 65–94 years). Central BP was 119.4 \pm 16.2 mmHg and 38.3 ± 11.6 mmHg, respectively for aortic systolic and pulse pressures. Mean pulse wave velocity (PWV) was 10.6 \pm 1.36 m/s and the augmentation index was 27.0 \pm 17.6%. Significant differences were depicted as a function of gender, with males presenting higher BP and PWV. The proportion of participants with increased PWV, according to the available reference values, was 31.6%. Participants with increased PWV had higher brachial and central BP, higher BMI and higher abdominal fat. Functionality was worst in high PWV participants, as well as cognitive function. Multivariate linear regression indicated age ($\beta = 0.172$; CI: 0.158;0.185; p < 0.001), and aortic systolic BP (β = 0.033; CI: 0.028;0.038; p < 0.001) as independent determinants of PWV. Also Hypertension (OR = 15.83; IC:8.16-30.7) and Diabetes (OR = 2.34; IC: 0.99-5.50) were independently associated with AS.

Conclusions: Accelerated AS is a common finding in the elderly and is highly associated with hypertension and diabetes. A strong association of AS with central BP and reflected waves is also of notice in this particular population.

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CORRELATION BETWEEN INFLAMMATORY STATE AND ARTERIAL STIFFNESS

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Pulse wave velocity (PWV) is gold standard for assessing arterial stiffness. Studies have shown that people with metabolic syndrome have insulin resistance and that after the onset of diabetes, cardiovascular risk is intensely increased, high-sensitive C-reactive protein (hsCRP) (1). Relate influence of changes in pulse wave velocity in the severity of the inflammatory state (2). Methods: A population-based cross-sectional study representative of a neighborhood of Salvador-BA, Brazil. The overall sample is randomized in adults from the assigned area, from December 2016 to May 2018 comprise 64 people. PWV was the measuring velocity between the carotid and right femoral wave. The flattening tonometer SphygmoCor® apparatus (XCEL, AtCor Medical, Australia). Blood samples were collected to biochemistry analysis, ADVIA1800[®] (SiemensHealthcare Japan/Canada). The committee for research FTC protocol (No1827621). Spearman's linear correlation coefficient between the laboratory tests and adjusted PWV were stratified according to the increased risk level of adjusted PWV. STATA v.12 for data analysis. The level of statistical significance was set at 5%.

Results: Table 1 (image 1), predominance of women (72.3%), (n = 64). When compared to the group with normal pulse wave velocity, there was an increase in the parameters of the laboratory tests in the group with an increased risk of arterial stiffness (adjusted PWV \geq 10), the correlations in this group and the PWV were positive and weak, except for the glycemia was negative, but they were not statistically significant. Already in the group with normal PWV, the correlations were positive and weak, only triglycerides presented. **Conclusion:** New molecular markers is necessary for correlate low intensity.

| Laboratory tests | Pulse wave velocity set ≥ 10 (n=17) | | | |
|------------------|-------------------------------------|-------------------|------------------------------|---------------|
| | Changed Parameters n (%) | (r; p-value) | Mean ± standard deviation | IC95% |
| Homa | 3 (17,7) | (r=0,0479;0,8553) | 3,0 ± 0,52 | 1,9 - 4,1 |
| hs-CRP | 6 (35,3) | (r=0,2611;0,3115) | 0,4 ± 0,1 | 0,2-0,6 |
| Triglycerides | 2 (11,8) | (r=0,1272;0,6225) | 149,2 ± 32,7 | 79,9 - 218,4 |
| Cholesterol | 6 (35,3) | (r=-0,0663;0,800) | 218,5 ± 12,1 | 192,9 - 244,1 |
| HDL | 14 (82,4) | (r=0,3434;0,1772) | 52,3 ± 2,6 | 46,8 - 57,8 |
| LDL | 7 (46.7) | (r=0,1243;0,6589) | 148,6 ± 10,5 | 126,2 - 171.0 |
| Insulin | 1 (5,9) | (r=0,0657;0,8022) | 11,2 ± 1,8 | 7,2 15,1 |
| Blood glucose | 8 (50,0) | (r=-0,0415;0,8787 | 112,9 ± 8,9 | 94,0 - 131,8 |
| | Pulse wave velocity set < 10 (n=47) | | | |
| Homa | 7 (14,9) | (r=0,1253;0,4012) | 2,6±0,3 | 2,0 - 3,3 |
| hs-CRP | 10 (21,3) | (r=0,1311;0,3799) | 0,3±0,1 | 0,2 - 0,5 |
| Triglycerides | 5 (10,6) | (r=0,3144;0,0314) | 110,5±9,9 | 90,6 - 130,4 |
| Cholesterol | 8 (17.0) | (r=0,2766;0,0599) | 199,1±5,9 | 187,2 - 211,0 |
| HDL | 40 (85,1) | (r=0,1559;0,2952) | 52,8±2,2 | 48.5 - 57.2 |
| LDL | 10 (21,3) | (r=0,1478;0,3216) | 125,7±5,1 | 115,5 - 135,9 |
| Insulin | 3 (6,4) | (r=0,0528;0,7245) | 10.4±1.2 | 8.1 - 12.7 |
| Blood glucose | 15 (31.9) | (r=0,2024;0,1725) | 99.3±3.6 | 92.0 - 06.7 |

References

 Mozos, Ioana et al. Inflammatory Markers for Arterial Stiffness in Cardiovascular Diseases. Frontiers in Immunology (2017) 8: 1058.
Bozkurt, B; Mann, D; Deswal, A. Biomarkers of inflammation in heart failure. Heart Fail Rev (2010) 15:331–341.

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ASSESSMENT OF CAROTID PULSE WAVE VELOCITY (CARPWV) IN PATIENTS WITH ANKYLOSING SPONDYLITIS

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