



Abstract 101 Figure 1 Survival of RVSP quartile.

**Conclusion** An RVSP of greater than 42 mm Hg is predictive of increased mortality in heart failure. This finding is independent of LVSD and COPD.

## 102 ETHNIC DIFFERENCES IN ENDOTHELIAL FUNCTION IN CHRONIC HEART FAILURE

doi:10.1136/heartjnl-2011-300198.102

<sup>1</sup>E Shantsila, <sup>2</sup>P S Gill, <sup>3</sup>G Y H Lip. <sup>1</sup>University of Birmingham Centre for Cardiovascular Sciences, City Hospital, Birmingham, UK; <sup>2</sup>University of Birmingham, Primary Care and Populational Sciences, Birmingham, UK; <sup>3</sup>University of Birmingham Centre for Cardiovascular Science, Birmingham, UK

**Background** Endothelial dysfunction is characteristic of patients with heart failure (HF) and is associated with an increased risk of future cardiovascular events. However, data on ethnic differences in endothelial function in HF are scarce. In this study we aimed to compare parameters of macro- and micro-vascular endothelial function and arterial elasticity in HF age- and sex-matched patients of different ethnic origin: (i) white European, (ii) south Asian and (iii) African-Caribbean. Additionally, SA patients with systolic HF were compared to two matched control groups: (i) south Asian patients with coronary artery disease without HF (disease controls) and (ii) south Asian “healthy controls”.

**Methods** We recruited 186 age/sex-matched patients with HF (ejection fraction <40%) of SA (n=43, age 66.5±11.1 years), white (n=44, age 68.4±9.4 years) and African-Caribbean (n=21, age 69.2±10.3 years) origin; as well as 36 disease controls (age 64.0±10.6 years) and 40 healthy controls (n=40, age 63.3±9.24 years). Macrovascular endothelial function was assessed as brachial artery flow mediated dilation in response to hyperaemia (FMD) and glyceryltrinitrate were assessed by vascular ultrasonography (iE33, Philips, USA). Microvascular endothelial function was evaluated by laser Doppler flowmetry of forearm skin (DRT4, Moor Instruments, UK) after iontophoresis of acetylcholine and sodium nitroprusside. Arterial stiffness was quantified by pulse wave velocity and augmentation index using (Sphygmocor, Australia).

**Results** Compared to disease controls and healthy controls south Asian patients with HF had impaired microvascular response to acetylcholine (390±302%, 549±264%, and 123±95.5%, respectively, p<0.05) and reduced FMD (7.12±3.64%, 11.8±4.66%, and 4.86±4.88%, respectively). HF patients of south Asian origin had impaired microvascular endothelial function (response to acetylcholine 123±95.5%) compared to white (258±15.6%) and African-Caribbean (286±17.3%) groups (p>0.05). HF patients of white origin had higher FMD than south Asian (4.86±4.88%) and African-Caribbean (5.36±3.24%) patients (p<0.05). No difference in

glyceryltrinitrate- and sodium nitroprusside-mediated (endothelial-independent) response was observed between study groups. In south Asian subjects, parameters of pulse wave velocity and augmentation index did not differ between those with HF and those in control groups. No ethnic differences were detected in pulse wave velocity. Conclusion: South Asian patients with HF have impaired micro- and macro-vascular endothelial function, but preserved arterial elastic properties. Significant ethnic differences in endothelial function are present in patients with HF.

## 103 SENILE SYSTEMIC AMYLOIDOSIS: A COMMON CAUSE OF HEART FAILURE IN THE ELDERLY?

doi:10.1136/heartjnl-2011-300198.103

<sup>1</sup>J H Pinney, <sup>2</sup>H J Lachmann, <sup>2</sup>J D Gillmore, <sup>2</sup>A Wechalekar, <sup>3</sup>S D J Gibbs, <sup>3</sup>P Sattianayagam, <sup>4,5</sup>S M Banypersad, <sup>6,7</sup>J Dzungu, <sup>3</sup>N Wassef, <sup>3</sup>C A McCarthy, <sup>3</sup>P N Hawkins, <sup>3</sup>C J Whelan. <sup>1</sup>National Amyloidosis Centre and UCL Centre for Nephrology, UCL Division of Medicine, Royal Free Hospital, London, UK; <sup>2</sup>National Amyloidosis Centre, UCL Division of Medicine, Royal Free Hospital, London, UK; <sup>3</sup>National Amyloidosis Centre, UCL Medical School, Royal Free Hospital, London, UK; <sup>4</sup>National Amyloidosis Centre, London, UK; <sup>5</sup>The Heart Hospital, UCL Medical School, London, UK; <sup>6</sup>National Amyloidosis Centre, UCL Medical School, University of London, London, UK; <sup>7</sup>St George's Hospital, University of London, London, UK

Senile systemic amyloidosis (SSA) is a rare cause of heart failure due to the deposition of wildtype transthyretin. The clinical features and outcome are ill defined; our aim was to evaluate the natural history of the disease in the UK in a group of thoroughly characterised patients. The series included all cases of biopsy proven transthyretin (TTR) amyloidosis with wildtype TTR gene sequencing who were prospectively followed up between January 2001 and May 2010. Clinical, biochemical, ECG and echocardiographic evaluation were performed at presentation to our centre. Patient survival was estimated using Kaplan–Meier analysis. 55 patients with histologically proven SSA; 36 (65.5%) from cardiac, 14 (25.4%) from GI tract, 3 (5.5%) from bladder, 1 (1.8%) from fat and 1 (1.8%) from carpal tunnel tissue were identified. 49 (89%) were male. The median age at diagnosis and death were 74 (range 66–89) and 79 (range 69–84) years respectively. Survival from symptom onset and diagnosis was 7.04 (range 0.54–8.41) and 4.58 (range 0.07–5.41) years respectively. In recent years more patients have been diagnosed with 2 (3.6%), 14 (25.5%) and 39 (70.9%) patients between 2001–2003, 2004–2006 and 2007–2009 respectively. The most common presentation was with breathlessness in 28 patients (51%). Twenty-four patients (43.6%) had prior carpal tunnel operations. Twelve (21.8%) patients had a history of ischaemic heart disease. Fifteen had had a coronary angiogram; 8 were reportedly normal and 7 required intervention. Arrhythmias were common, 20 patients (36.3%) had a history of atrial fibrillation and 6 (10.9%) had pacemakers in situ. ECG findings were; 24 (43.6%) in AF, 6 (10.9%) first degree block, 10 (18.2%) left bundle and 6 (10.9%) right bundle branch block, 27 (49%) T wave changes, 11 (20%) <5 mm complexes in all inferior leads. Echocardiographic findings revealed the median IVSd was 1.7 (range 1.1–2.5) cm, median E/A ratio was 2.7 (range 0.79–5.4), E/E' 15.81 (range 7.5–41.1) and ejection fraction was 45.5 (range 13–83)%. Blood results showed; the median baseline NT-proBNP was 356.1 (range 5–2611) and troponin T 0.03 (range 0.01–0.28). Twenty-five patients had a troponin T >0.03 (45%). Ten patients (18%) had a detectable paraprotein and 2 (3.6%) had bence jones proteins. SSA is present in >25% of the very elderly at post mortem but was rarely diagnosed during life. It is becoming more frequently recognised perhaps due to widespread use of cardiac MRI. Most patients are male but women can be affected. A history of carpal tunnel syndrome is common. The diagnosis is often made after the onset of breathlessness. Systolic and diastolic dysfunction