





Assessing the reproducibility of aortic stiffness in patients with end-stage renal disease: a 3-Tesla cardiac magnetic resonance imaging study

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Background

Cardiovascular disease is the leading cause of death in patients with endstage renal disease (ESRD) but is driven by a different set of risk-factors than classical coronary artery disease. One of these risk factors is increased aortic stiffness, which disrupts arterio-ventricular coupling and leads to left ventricular hypertrophy. Aortic stiffness is an independent predictor of cardiovascular mortality in ESRD and a potential imaging biomarker. It is measurable with cardiac magnetic resonance (CMR) imaging using:

Results

	Population (n=10			
Male; n (%)	8 (80)	Table 1: Patient demographics		
Age; years (mean ± SD)	59 ± 15	Normally distributed data		
HD vintage (months)	26.2 (±26.6)	presented as mean (±SD); non-		
Active on transplant list; n (%)	5 (50)	normally distributed data as median (25 th , 75 th percentile) BAME, Black, Asian and		
Ethnicity; n (%)				
Caucasian	3 (30)	Minority Ethnic		
BAME	7 (70)			

- Aortic pulse wave velocity (aPWV) is an indirect measure of aortic stiffness
- Aortic distensibility (AD), a direct measure

For the first time in this population, the inter-study repeatability and the intraand inter-observer variability of aPWV and AD was investigated with high resolution 3-Tesla CMR.

Methods

10 haemodialysis patients underwent two identical test-retest CMR scans within 2 weeks.

Analysis was offline and blinded. Aortic PWV was calculated from saggitaloblique cine images of the aortic arch to measure distance (Figure 1a) and phase contrast sequences of the ascending and descending aorta to derive transit time (Figure 1b-c). Ascending and descending AD (AAD and DAD) was calculated from axial cine images, at the level of the pulmonary artery bifurcation (Figure 1d).

All 10 patients completed the study. Demographics are presented in Table 1. Mean aPWV was 8.2m/s (±3.48), mean AAD was 2.5mmHg⁻¹x10⁻³ (±1.7), mean DAD was 3.2mmHg⁻¹x10⁻³ (±2.7). The repeatability and variability of parameters is shown (Table 2). Bland-Altman analyses revealed no systematic bias (Figure 2a-c).

Table 2: Reproducibility of aPWV, ascending AD and descending AD.

* Three inter-study aPWV scans were not analysable and excluded from analysis

	aPWV (m/s) (n=7)	AAD (mmHg ⁻¹ x10 ⁻³) (n=10)	DAD (mmHg ⁻¹ x10 ⁻³) (n=10)
Inter-study repeatability	ICC: 0.51 (-0.31, -0.90)* CoV: 14.8%	ICC: 0.94 (0.78, 0.99) CoV: 13.3%	ICC: 0.51 (-0.13, 0.85) CoV: 33.7%
Inter-observer variability	ICC: 1.00 (1.00,1.00) CoV: 1.4%	ICC: 1.00 (0.99, 1.00) CoV: 2.7%	ICC: 1.00 (0.99 <i>,</i> 1.00) CoV: 2.1%



Inter-study repeatability was assessed by a single reader. Inter- and intraobserver variability were respectively assessed by two readers of 10 scans and one reader analysing 10 scans twice. Intra-class correlation coefficient (ICC), coefficient of variation (CoV) and Bland-Altman analyses were calculated to assess reproducibility.





Figure 2: Inter-study repeatability: a) aPWV; b) AAD; c) DAD; table shows bias and *limits of agreement.*

Figure 1: *CMR* analysis of aPWV and AD.

Conclusions

Inter-study results show that ascending AD is the most reproducible measure of a stiffness. Furthermore, the ascending a statistic potentially the most clinically relevant area for arterio-ventricular coupling in ESRD, as it has been shown to:

- Have the most capacitance
- Be most affected by aortic stiffening \bullet
- Most strongly associate with increased LV mass

The inter- and intra-observer variability for all parameters were excellent. The inter-study repeatability of aPWV and descending AD suggest they are suboptimal for use as biomarkers of aortic stiffness.

