

POSTER PRESENTATION

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Predicting hemodynamics in native and residual coarctation: preliminary results of a Rigid-Wall Computational-Fluid-Dynamics model (RW-CFD) validated against clinically invasive pressure measures at rest and during pharmacological stress

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From 2011 SCMR/Euro CMR Joint Scientific Sessions
Nice, France. 3-6 February 2011

Introduction

Current clinical assessment of borderline aortic coarctation may involve cardiovascular magnetic resonance (CMR) but if inconclusive, invasive catheterization pressure measurements are required to evaluate the pressure gradient at rest and during pharmacological stress (isoprenaline).

Purpose

To predict the aortic pressure distribution in patients with aortic coarctation at rest and pharmacological stress using a transient rigid-walled computation fluid dynamics model(RW-CFD).

Methods

The study cohort comprises 5 patients with native or recurrent aortic coarctation and 2 control patients with healthy aortic arches(Table 1), who underwent both CMR(1.5-Intera,Philips) and catheterization at rest and pharmacological stress.

The model workflow(Figure 1) requires,as input parameters,the aortic geometry, extracted from the CMR 3D gadolinium contrast-enhanced sequence(TR=4.4ms, TE=1.8ms,1.5x1.5x1.8mm), and definition of the

boundary conditions. The blood flow(modelled as a Newtonian incompressible fluid) in the aortic domain is conditioned by the clinical data at three locations: **1) Ascending aortic root:** The inlet flow is extracted from the phase-contrast CMR flow(TR=4.7ms,TE=3ms, 2.5x2.5x7mm,80 phases). **2) Supra-aortic vessels:**The flow rate is calculated as a proportion of the inlet flow based on the assumption of a constant wall shear stress (Kundu,2004).**3) Diaphragmatic aorta:** The pressure waveform is extracted from the invasive catheter investigation.

The clinically invasive aortic pressure gradients were compared with the predicted pressure distribution along the centreline in the RW-CFD model at the time of peak flow(Table 2).

Results

For patients with aortic coarctation, during pharmacological stress, there was an increase in both heart rate (72 ± 21 bpm, *mean \pm standard deviation*) and invasive pressure gradient drop across the coarctation (35 ± 18 mmHg, Table 2). The RW-CFD model predicted accurately the pressure drop at rest (-4.2 ± 4.9 mmHg), and moderate agreement at stress (-4.4 ± 21.9 mmHg. Table 2, Figure 3).

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Table 1 Study population demographics

Study number	Age [years]	Weight [kg]	Body surface area [m ²]	Clinical condition	Previous procedures
Aortic coarctation					
AoCo-1	15	59	1.7	Residual aortic coarctation, mitral stenosis	Coarctation repair (end-to-end anastomosis)
AoCo-2	25	64	1.8	Residual aortic coarctation, atrial septal defect	Coarctation repair (end-to-end anastomosis) and ASD repair
AoCo-3	21	95	2.1	Residual aortic coarctation, bicuspid aortic valve	Coarctation repair (Dacron patch)
AoCo-4	20	71	1.9	Residual aortic coarctation	Coarctation repair (subclavian flap)
AoCo-5	17	71	1.9	Native aortic coarctation	None
Normal aorta					
1	2	15	0.6	Partial anomalous pulmonary venous return, residual pulmonary vein stenosis	PAPVR repair
2	1	8.4	0.4	Biliar artesia	None

For healthy controls, the RW-CFD model predicted the absence of a significant gradient both at rest and stress (1 ± 1 mmHg).

Conclusion

For patients with aortic coarctation, the RW-CFD simulations accurately predict the pressure gradient at rest and give indication of the gradient severity during stress. Furthermore, no gradient was predicted in control patients with normal aortae.

These preliminary results, whilst using a simple CFD approach and a small cohort of patients, are quite promising. This study represents the first step towards an image-based fluid-solid-interaction CFD analysis. This more sophisticated approach is likely to overcome the current limitations and might grant additional information.

In the future, it is envisaged that CFD models could be based on a patient-specific, non-invasive and non-ionising radiation assessment such as CMR in order to

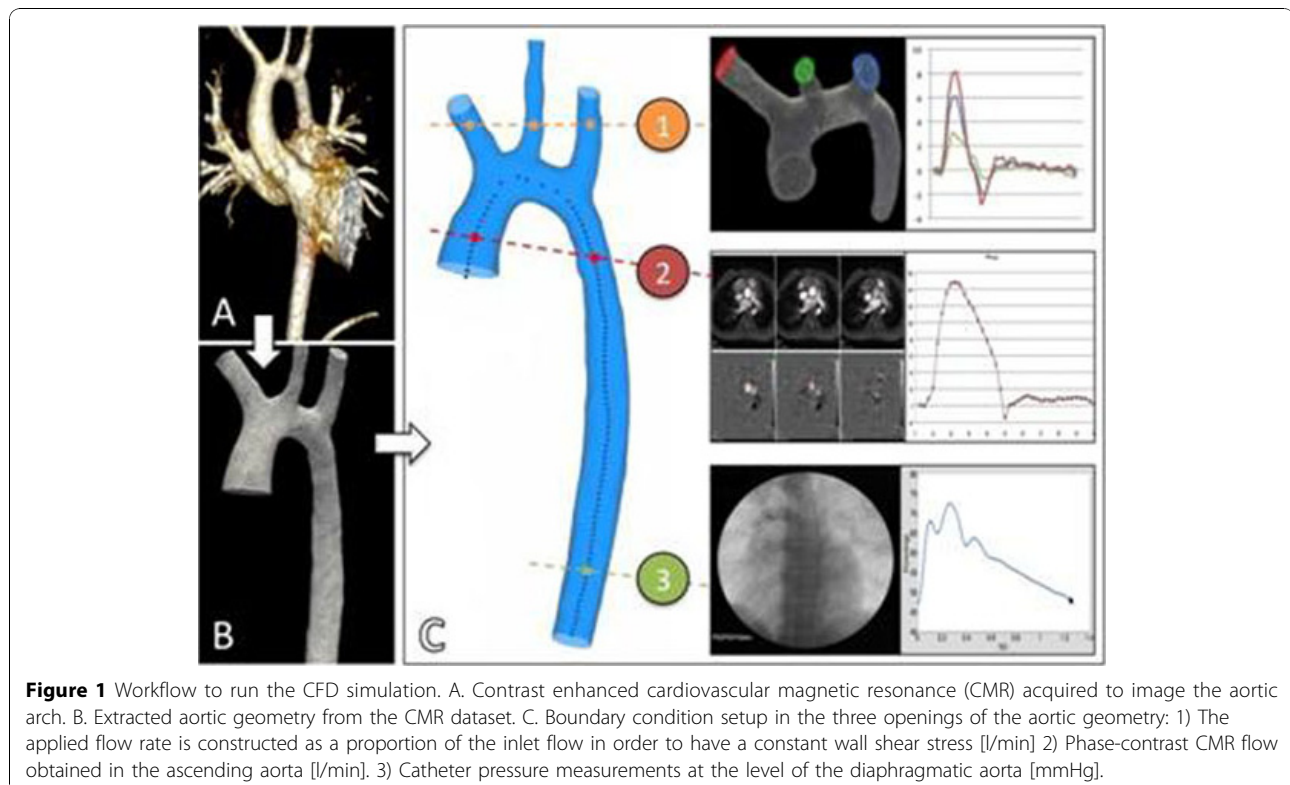
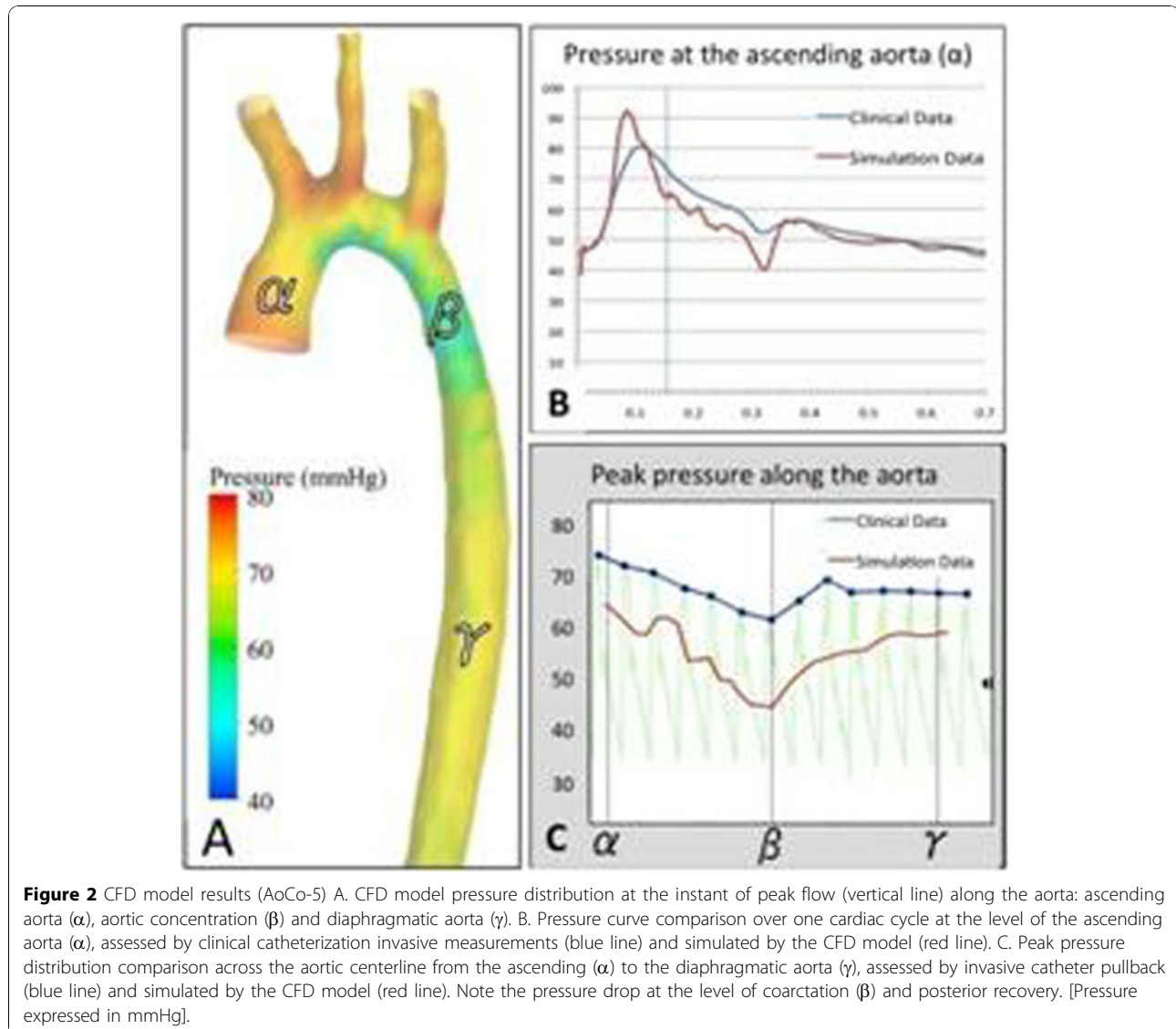


Table 2 Study results in patients with aortic coarctation (AoCo) at rest and stress conditions

Study number	Heart rate [bpm]	Cardiac Output [l/min/m ²]	Pressure Ascending Aorta Invasive [mmHg]	Pressure Diaphragmatic Aorta Invasive [mmHg]	ΔP Clinical Invasive [mmHg]	ΔP CFD [mmHg]	Pressure Difference (CFD - Invasive) [mmHg]
Rest condition							
AoCo-1	48	2.0	102 ± 3	79	23 ± 3	22	-1 ± 3
AoCo-2	86	3.2	82 ± 3	64	18 ± 3	5	-13 ± 3
AoCo-3	69	2.2	86 ± 4	74	12 ± 4	9	-2 ± 4
AoCo-4	81	2.8	78 ± 2	68	10 ± 2	8	-2 ± 2
AoCo-5	47	1.9	82 ± 2	73	9 ± 2	6	-3 ± 2
Stress							
AoCo-1	150	5.6	116 ± 6	77	39 ± 6	54	18 ± 6
AoCo-2	136	5.4	103 ± 10	63	40 ± 10	23	-17 ± 10
AoCo-3	130	5.6	121 ± 6	57	64 ± 6	42	-22 ± 6
AoCo-4	140	6.5	152 ± 4	86	66 ± 4	44	-22 ± 4
AoCo-5	141	7.2	114 ± 7	77	37 ± 7	58	21 ± 7



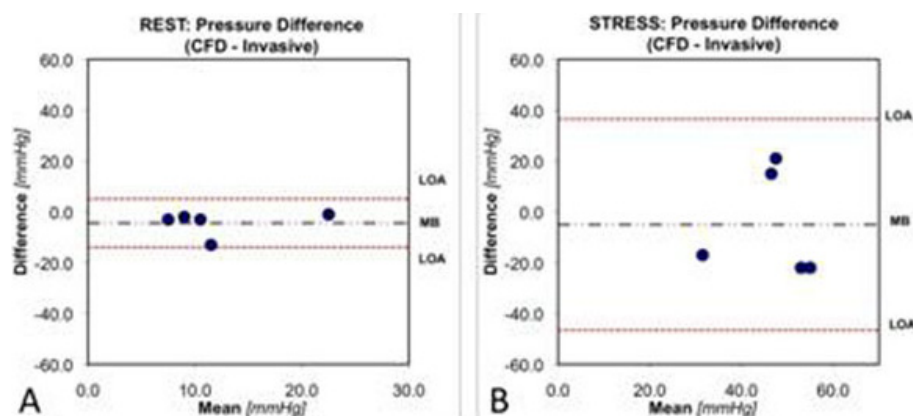


Figure 3 Bland-Altman plots: Comparison of the clinically invasive and the predicted CFD pressure gradient (mmHg) across the aortic coarctation at rest (A) and isoprenaline pharmacological stress (B). The dot-dashed grey horizontal lines represent the mean difference between CFD and invasive data (MB, mean bias), and the paired dotted red horizontal lines represent ± 2 standard deviations from this mean difference (LOA, 95% limits of agreement). Note the increased bias during the stress condition compared to the good agreement at rest condition.

predict the hemodynamic conditions in the aorta and avoid invasive cardiac catheterization.

The pressures assessed by clinically invasive catheterization at the level of the ascending aorta, diaphragmatic aorta and the pressure gradient across the coarctation (ΔP Clinical Invasive) are compared with the RW-CFD model pressure gradient prediction (ΔP CFD). The absolute pressure differences values are shown in the final column. Pressures expressed as mean \pm standard deviation. bpm, beats per minute.

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Published: 2 February 2011

doi:10.1186/1532-429X-13-S1-P49

Cite this article as: Valverde *et al.*: Predicting hemodynamics in native and residual coarctation: preliminary results of a Rigid-Wall Computational-Fluid-Dynamics model (RW-CFD) validated against clinically invasive pressure measures at rest and during pharmacological stress. *Journal of Cardiovascular Magnetic Resonance* 2011 **13**(Suppl 1):P49.

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