

# **POSTER PRESENTATION**

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# MR-augmented cardio pulmonary exercise testing: an integrated approach to assessment of children with pulmonary hypertension

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## **Background**

Pulmonary arterial hypertension (PAH) is a life limiting disorder resulting in progressive right heart (RV) failure. Cardiac MRI is the reference standard for RV assessment in PH and has known prognostic value in children. Symptoms of PAH are exacerbated or become apparent with exercise and cardiopulmonary exercise testing (CPET) is also prognostic in PAH.

We have recently demonstrated the feasibility of MR augmented CPET (MR-CPET) in healthy adults. Using the Fick principle MR-CPET allows calculation of tissue oxygen extraction  $\Delta cO_2$ . A combined approach using MR-CPET could provide a more sensitive method of assessing disease severity, the causes of exercise limitation and response to treatment. In this study we assessed the utility of MR-CPET in children with PAH.

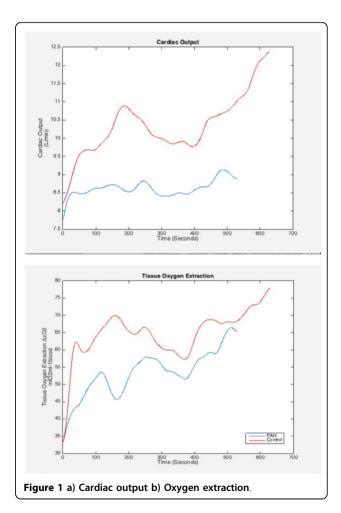
#### Methods

10 Children with PAH aged 8 to 16 years (WHO class I or II, on oral/inhaled therapy only) and 10 healthy volunteers matched for age and sex underwent MR-CPET. Exercise was performed on a MRI compatible ergometer (Lode, Groningen, The Netherlands) CPET data was collected using a commercial respiratory gasanalyser (Medgraphics, St. Paul, USA) with a modified MR compatible sampling umbilical.

Resting and peak exercise ventricular volumes and septal curvature were measured using a radial k-t SENSE real time sequence. Aortic flow was simultaneously continuously measured using a previously validated real-time UNFOLD-SENSE spiral PCMR sequence.

MR data was used to derive CO, HR and SV curves during exercise. Arteriovenous oxygen content gradient

(AVO2) curves (a measure of tissue oxygen extraction) were calculated by dividing the VO2 and CO curves (Figure 1). Resting and peak exercise mean pulmonary



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artery pressures (mPAP) were estimated based on septal curvature using a previously validated technique.

Subjects also completed a six-minute walk test, experience survey and exercise history.

### **Results**

All children successfully completed the test without complication. Peak  $VO_2$  was significantly lower in the PAH

group. This seems to be driven by both reduced tissue oxygen extraction and cardiac output although only reduced  $\Delta cO_2$  was of statistical significance. MR-CPET measures at rest and exercise are shown in Figure 2.

RVEF was significantly lower in the PAH group both at rest and at peak exercise. There was a significant difference in the increase in estimated mPAP between the control and PAH groups. There was no significant

Variable	Control group	PAH group	p Value
Resting VO <sub>2</sub> (ml/min <sup>-1</sup> )	191 (157- 232)	181 (161- 204)	0.6182
Peak VO <sub>2</sub> (ml/min <sup>-1</sup> )	803 (676- 931)	578 (463- 694)	.00834
Resting cardiac output, Lmin <sup>-1</sup>	6.5 (5- 8)	6.1 (5.3-7)	.6359
Peak cardiac output, Lmin <sup>-1</sup>	9.8 (8-12)	8.1 (7- 9.2)	.0831
Resting stroke volume, ml/beat	71 (56-85)	66 (57- 76)	.5742
Peak Stroke volume, ml/beat	67 (54-80)	62 (51-72)	.502
Resting heart rate, bpm	83 (71-95)	81 (74-88)	.7924
Peak heart rate, bpm	149 ( 136- 161)	133 (121- 145)	.06275
Resting tissue extraction $\Delta cO_2$ , mIO <sub>2</sub> mI <sup>-1</sup> blood	191 ( 157- 232)	181 (61- 204)	0.6182
Peak tissue extraction ΔcO <sub>2</sub> , mlO <sub>2</sub> ml <sup>-1</sup> blood	803 (676- 931)	578 (463- 694)	.00834
VE/VCO <sub>2</sub>	38 (35-42)	43 (39-47)	.04454
Estimated mPAP rest, mmHg	11 (9- 14)	32 (25- 41)	0.00
Estimated mPAP exercise, mmHg	15 (12-17)	58 (47- 71)	0.00
Difference mPAP, rest – exercise. mmHg	3.7 (2.2-6.1)	20 (12- 35)	0.0001
RVEF rest	62 (58-65)	56 (51-60)	.03002
RVEF peak	69 (63-75)	60 (55-65)	.01855
Six minute walk test distance, m	452 (418- 489)	451 (400-507)	0.9578

Figure 2 MR-CPET findings.

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difference in 6-minute walk test distance between the groups.

#### **Conclusions**

MR-CPET is well tolerated in both healthy children and children with class I/II PAH. This study demonstrates that exercise intolerance in children with PAH is in part due to reduced tissue oxygen extraction and may open possibilities to study new pharmacological or exercise based therapeutic strategies.

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