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# 1000 Ventricular-vascular coupling is independently associated with exercise capacity in patients with ischemic cardiomyopathy: a cardiac multi-modality imaging study

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### Introduction

Left ventricular (LV) systolic dysfunction due to ischemic cardiomyopathy (I-CMP) leads to reduced exercise capacity. Ventricular-vascular coupling (VVC), a ratio of effective arterial to LV elastance, represents forward flow efficiency of LV, independent of mitral regurgitation.

### **Purpose**

In I-CMP patients, we sought to a) assess relationship between resting VVC and maximum oxygen consumption corrected for peak heart rate (MVO<sub>2</sub>/pHR), an accurate measure of exercise capacity in patients on  $\beta$ -blockers, and b) compare value of VVC versus other determinants of exercise capacity.

#### Methods

43 patients with I-CMP (age 59  $\pm$  9 years, 88% on  $\beta$ blocker) underwent cardiopulmonary exercise testing, echocardiography and cardiac magnetic resonance (CMR, 1.5 T Siemens Scanners, Erlangen Germany) for cardiac transplant evaluation. MVO<sub>2</sub>/pHR and diastolic filling variables (echocardiography) were measured in a standard fashion. CMR LV indices [end-systolic (ESV), enddiastolic (EDV), stroke volume (SV), all in ml, and LVEF were measured using the standard contiguous short-axis slices from apex to base, using the balanced steady state free precession cine sequence (TE = 1.6 msec, TR = 3.3 msec, flip angle = 70° and slice thickness 8-10 mm, field of view varied from 228-330 in the x-direction and 260-

330 in the y-direction and matrix size varied from 140-180 in the x-direction and 256 in the y-direction, giving a spatial resolution of 1.5-2.1 mm (x-direction) by 1.1-1.4 mm (y-direction). For patients able to suspend respiration, breath hold duration was 10-15 sec, depending on the heart rate; otherwise, images were acquired using 3 signal averages. Subsequently, off-line analysis was performed using Argus analytical software (Siemens Medical Solutions, Erlangen, Germany) to assess LV volumes and LVEF, in a standard fashion. VVC was calculated as: [Effective arterial elastance (end systolic pressure ÷ stroke volume index)/LV end-systolic elastance (end systolic pressure ÷ LV systolic volume index)].

## Results

Mean LVEF, ESV, EDV, and SV were 24 ± 8%, 228 ± 107 ml, 297  $\pm$  111 ml, and 66  $\pm$  20 ml respectively. Mean  $MVO_2/pHR$  was 13 ± 3 ml/beat. Predictors of  $MVO_2$  are shown in Table 1. On stepwise regression, only VVC significantly predicted MVO<sub>2</sub>/pHR.

#### Conclusion

In I-CMP patients with LV dysfunction, VVC predicts MVO<sub>2</sub>/pHR that can potentially be used as a potential therapeutic target.

Table I: Univariate and multivariate predictors pf MVO<sub>2</sub>/peak HR.

	MVO <sub>2</sub> /peak HR		
	Correlation coefficient (rho)	P value	Multivariate analysis P value
Clinical characteristics			
Age	-0.006	0.9	
Gender	0.1	0.5	
NYHA class	-0.14	0.4	
Baseline SBP	0.22	0.16	
Baseline DBP	-0.02	0.9	
Echocardiographic and CMR features			
E max	-0.39	0.009	0.1
E/A ratio	-0.23	0.1	
E wave deceleration time	0.35	0.02	0.5
CMR features			
LV end-systolic volume	-0.41	0.006	0.9
LV end-diastolic volume	-0.28	0.07	
Stroke volume	0.14	0.3	
LV ejection fraction	0.36	0.02	0.6
Ventricular vascular coupling	0.45	0.002	0.001
Ea .	-0.28	0.09	
Ees	0.39	0.009	

NYHA indicates New York Heart Association; SBP, systolic blood pressure; DBP, diastolic blood pressure; E max, mitral E wave velocity; LV, left ventricle; Ea, Arterial elastance; Ees, left ventricular systolic elastance.

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