Abstract 26 Figure 1 Panel A: scatter plot of measured and predicted mean pulmonary artery pressure. Panel B: Main pulmonary artery (MPA) time-resolved, flow curves (the light blue area was used to compute mean pan-systolic area)

Abstract 26 Table 1 Summary of CMR parameters evaluated in the study

	Male (n=8)		Female (n=10)		P-value
	Mean	SD	Mean	SD	
Age (Years)	71.0	6.8	66.0	7.8	0.16
Invasive mPAP (mmHg)	36.4	9.5	41.9	8.9	0.23
RA Area (cm ²)	23.7	6.1	28.1	11.8	0.31
RVEDV (ml)	165.3	51.1	202.3	72.6	0.22
RVESV (ml)	114.8	49.1	138.0	58.7	0.37
RVSF (%)	50.7	25.5	64.4	22.7	0.25
RVEF (%)	32.5	15.7	33.2	8.9	0.91
PA SV (ml)	50.0	24.7	53.6	16.8	0.72
PA Mean Systolic Velocity (cm/s)	15.0	6.9	14.3	3.4	0.79
PA WSR	39.5	22.1	36.6	10.2	0.74
PA WSS	138.1	77.3	127.9	35.2	0.73
Predicted mPAP CMR MODEL (mmHg)	37.5	7.4	40.5	6.8	0.40

acquisition. The velocity encoded images were analysed for the following: mean pulmonary artery pan-systolic velocity (PASV), MPA stroke volume, MPA wall shear stress (WSS) and wall shear rate (WSR). The 4-chamber cine was used to measure end-diastolic right atrial area. Right ventricular volumes were analysed using standard methods. Stepwise multiple regression model of significantly associated parameters ($p < 0.05$) was developed.

Results Mean age of the 18 patients was 68.78 ± 7.46 years (44% males). The following CMR metrics demonstrated significant association to the measured mPAP: RA area ($r = 0.65$ $p = 0.03$); MPA mean pan-systolic velocity ($r = -0.57$ $p = 0.01$); RVEDV ($r = 0.52$ $p = 0.03$); RVEF ($r = -0.40$ $p = 0.10$); RVESV ($r = 0.58$ $p = 0.01$). In stepwise multiple regression, only two parameters demonstrated independent association to mPAP - RA Area and MPA mean pan-systolic velocity. The predicted mPAP demonstrated good correlation to the measured mPAP ($R = 0.76$, $p < 0.001$).

Conclusion(s) Mean pulmonary artery systolic velocity and right atrial area are independently associated with mPAP.

Our novel CMR prediction model for mPAP, comprising of these two metrics, demonstrates high association to the measured mPAP by invasive haemodynamic study.

27 MIXED VENOUS OXYGEN LEVELS IN PULMONARY HYPERTENSION IS ASSOCIATED WITH RIGHT HEART MULTI-PARAMETRIC ASSESSMENT

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Background Pulmonary hypertension (PH) is debilitating disease characterized by a progressive increase in pulmonary arterial pressure (PAP) that leads to right ventricular (RV) failure and death. Mixed venous oxygen saturation (SVO₂) represents the oxygen saturation of blood returning to the lungs before reaching the alveolo-capillary units. SVO₂ is strongly associated with clinical outcomes in PH. The relationship of non-invasive CMR metrics to this prognostically relevant parameter in patients with PH are unknown.

Purpose This study sought to develop an early understanding of which CMR volumetric and flow parameters are most associated with SVO₂.

Methods Eighteen ($n = 18$) patients were prospectively recruited at a large tertiary PH unit. The SVO₂ was measured during right heart catheterisation. All patients had CMR on the 1.5 T scanner (HDx scanner, GE Healthcare, Waukesha, Wisconsin, USA), using an 8-channel cardiac coil. Subjects were scanned in the supine position with electrocardiogram (ECG) gating. CMR protocol included long and short axis cines and through-plane pulmonary artery phase contrast acquisition. The velocity encoded images were analysed for the following: mean pulmonary artery (MPA) systolic velocity, MPA stroke volume, MPA wall shear stress (WSS) and wall shear rate (WSR). The 4 chamber cine was used to measure end-diastolic right atrial (RA) area. RV volumes were analysed using standard methods. Stepwise multiple regression model of significantly associated parameters ($p < 0.05$) was developed.

Abstract 27 Figure 1 Panel A: scatter plot of measured and predicted SVO2. Panel B: Right atrial size was computed in 4-chamber view at end-systole (RV), just before the opening of the tricuspid valve

Abstract 27 Table 1 Summary of CMR parameters evaluated in the study

CMR parameters	Mean	SD	Pearson Correlation (P-value)	Regression model*
RA Area* (cm2)	25.64	9.05	−0.57 (0.01)	F-ratio=8.56
PA SV* (mls)	51.58	21.05	0.5 (0.03)	p<0.01
LVEDV (mls)	119	39.62	0.42 (0.08)	
LVESV (mls)	54.92	16.57	0.22 (0.38)	
LVSV (mls)	63.39	29.97	0.42 (0.08)	
LV mass (grams)	82.61	22.03	0.27 (0.3)	
LVEF (%)	53.02	10.34	0.26 (0.29)	
RVEDV (mls)	181.72	62.52	−0.12 (0.63)	
RVESV (mls)	125.08	53.26	−0.30 (0.22)	
RVSV (mls)	56.78	24.59	0.35 (0.16)	
RVEF (%)	32.81	12.79	0.46 (0.05)	

Results The left ventricular and RV volumetric functional parameters demonstrated no association to SVO2 ($p>0.05$) in any of the participants. However, a negative association was shown between RA area and SVO2 ($R=-0.57$, $p=0.01$). The only other parameter which correlated with SVO2 was MPA stroke volume ($R=0.5$, $p=0.03$). In stepwise multiple regression, both parameters demonstrated independent association to SVO2. The predictive values generated by a combined model demonstrated high correlation to measured SVO2 ($R=0.73$, $p<0.001$).

Conclusion(s) RA area and MPA stroke volume are independently associated with SVO2. A novel CMR prediction model comprised of these two metrics, demonstrates high association to the measured SVO2 by invasive haemodynamic study. Further work to determine reproducibility and reliability is needed before this method becomes an important prognostic tool for PH patients.