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Repeatability and sensitivity to change of non-invasive end points in PAH: the RESPIRE study

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Abstract End points that are repeatable and sensitive to change are important in pulmonary arterial hypertension (PAH) for clinical practice and trials of new therapies. In 42 patients with PAH, test–retest repeatability was assessed using the intraclass correlation coefficient and treatment effect size using Cohen’s d statistic. Intraclass correlation coefficients demonstrated excellent repeatability for MRI, 6 min walk test and log to base 10 N-terminal pro-brain natriuretic peptide (\log_{10} NT-proBNP). The treatment effect size for MRI-derived right ventricular ejection fraction was large (Cohen’s d 0.81), whereas the effect size for the 6 min walk test (Cohen’s d 0.22) and \log_{10} NT-proBNP (Cohen’s d 0.20) were fair. This study supports further evaluation of MRI as a non-invasive end point for clinical assessment and PAH therapy trials.

Trial registration number NCT03841344.

INTRODUCTION

Pulmonary arterial hypertension (PAH) is progressive, leading to right ventricular (RV) failure and death.¹ Accurate measurement of RV function is important for assessment of disease severity and prognosis.^{2–4} Despite new therapies and improvements in survival,⁵ PAH remains a life-shortening condition. MRI is the gold standard for RV assessment,⁶ has prognostic value² and predicts clinical worsening⁷ in PAH. A trial end point that is highly repeatable, is sensitive to treatment and predicts outcomes would be highly desirable.^{8,9} MRI has been proposed as a trial end point in PAH,^{8,9} however, there is limited data on repeatability and treatment effect size.

METHODS

Patients

Patients with PAH who were treatment-naïve commencing therapy, prevalent undergoing escalation of therapy and clinically stable requiring no escalation of therapy, were recruited. See online supplemental file S1.

Study investigations

Investigations performed at visit 1 included N-terminal pro-brain natriuretic peptide (NT-ProBNP), 6 min walk test (6MWT) and MRI. Follow-up visits 2 and 3 occurred approximately 6 months after study visit 1. Visits 2 and 3 occurred within 24 hours of each other (online supplemental figure S2).

MRI acquisition and analysis

All MRI examinations were performed on either a 1.5 T GE HDx (GE Healthcare, Milwaukee, USA) whole body scanner using an 8-channel cardiac coil or a 3 T Philips Ingenia (Best, The Netherlands) whole body scanner using a 32-channel dStream torso coil (online supplemental file S1). Analysis of MRI was undertaken blinded to the patient’s data. RV parameters and pulmonary arterial flow were analysed on Qmass MEDIS suite (V3.0.18.0, Medical Imaging Systems, The Netherlands) on short axis and phase contrast images, respectively. Regions of interest were drawn on the pulmonary artery and left atrium of the dynamic contrast-enhanced perfusion images to calculate first pass pulmonary transit time and full width at half maximum using in-house software (see online supplemental figure S3).

Six min walk test and NT-ProBNP

The 6MWT was performed by a respiratory physiologist. NT-ProBNP analysis was performed on patient plasma samples using the Luminex 100/200 multiplex analyser using the cardiovascular marker kit (HCVD-1MAG-67K Millipore) at the end of the study.

Statistical analysis

Repeatability was determined by the intraclass correlation coefficient (ICC) using a two-way mixed absolute agreement model with the average measure recorded. An ICC of ≥ 0.75 was considered excellent, 0.60–0.74 good, 0.40–0.59 fair and < 0.40 poor. Mean difference and 95% CIs were presented where appropriate. Cohen’s d (calculated with the averaged SD, d_{av}) was used to assess the standardised treatment effect size between visit 1 and visit 2.¹⁰ A Cohen’s d value of < 0.20 was considered no change, 0.20–0.49 was considered fair change, 0.50–0.79 was considered a medium change and ≥ 0.80 was considered a large change. All analysis was performed on SPSS V.22 and GraphPad Prism V.16.

RESULTS

Patients

Of 42 patients who completed the study, 16 were incident and treatment-naïve and initiated PAH therapy, 12 were prevalent and underwent an escalation of therapy and 14 were stable on therapy with no change in treatment occurring between the study visits. (online supplemental table S5).



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Table 1 Repeatability in all patients with PAH (ICC), and treatment effect size for patients with PAH initiating or escalating PAH therapy

	Patients with PAH initiating or escalating therapy												
	All PAH			Visit 1		Visit 2		Change (Visit 1–visit 2)		95% CI			
	N	ICC	N	Mean	SD	Mean	SD	Mean difference	SD	SEM	Lower	Upper	Cohen's d
Walk test													
6MWT distance (m)	39	0.987	24	325.63	156.30	361.50	166.29	-35.88	79.06	16.14	-69.26	-2.49	0.22
Blood tests													
Log NT-ProBNP	32	0.772	24	2.76	0.46	2.67	0.41	0.09	0.32	0.07	-0.05	0.22	0.20
MRI metrics													
<i>SA with threshold</i>													
RVEDM (g)	40	0.970	26	117.80	45.72	99.40	43.96	18.40	30.90	6.06	5.92	30.88	0.41
RVESM (g)	40	0.980	26	106.68	39.73	94.61	42.08	12.06	26.79	5.25	1.24	22.88	0.29
RVEDV (mL)	40	0.969	26	145.71	39.12	146.03	55.87	-0.32	29.13	5.71	-12.08	11.45	0.01
RVESV (mL)	40	0.983	26	93.93	34.66	81.28	41.40	12.65	22.02	4.32	3.76	21.55	0.33
RVEF (%)	40	0.883	26	36.56	11.48	45.69	11.12	-9.12	10.45	2.05	-13.35	-4.90	0.81
RVSV (mL)	40	0.864	26	51.78	17.30	64.75	23.92	-12.97	23.27	4.56	-22.37	-3.57	0.62
RVCO (L/min)	40	0.886	26	3.95	1.45	4.48	1.55	-0.53	1.54	0.30	-1.15	0.09	0.35
Systolic septal angle (°)	40	0.852	27	163.33	16.45	156.81	14.00	6.52	11.28	2.17	2.06	10.98	0.43
Diastolic septal angle (°)	40	0.897	27	153.11	14.73	145.48	10.44	7.63	10.15	1.95	3.61	11.65	0.60
Q flow													
Net flow volume (mL)	41	0.893	26	58.05	30.18	69.49	31.30	-11.44	34.83	6.83	-25.51	2.62	0.37
Forward flow volume (mL)	41	0.860	26	60.37	27.58	72.33	29.15	-11.96	31.97	6.27	-24.88	0.95	0.42
Backward flow volume (mL)	41	0.817	26	2.32	6.76	2.84	5.74	-0.52	5.85	1.15	-2.88	1.85	0.08
Regurgitant fraction (%)	41	0.731	26	6.28	19.58	5.42	11.52	0.87	18.77	3.68	-6.71	8.45	0.05
Average flow velocity (cm/s)	41	0.909	26	7.31	3.60	8.25	3.69	-0.94	3.69	0.72	-2.43	0.55	0.26
Peak flow velocity (cm/s)	41	0.582	26	52.97	16.37	67.68	22.71	-14.71	19.35	3.79	-22.53	-6.90	0.74
Diastolic vessel area (mm ²)	41	0.933	26	981.10	257.92	961.84	242.96	19.26	104.52	20.5	-22.96	61.48	0.08
Systolic vessel area (mm ²)	41	0.953	26	1077.57	279.96	1083.62	266.78	-6.05	101.08	19.82	-46.88	34.77	0.02
Pulmonary arterial pulsatility (%)	41	0.776	26	9.96	4.87	13.00	5.12	-3.04	3.62	0.71	-4.50	-1.58	0.61
DCE imaging													
Pulmonary transit time (s)	36	0.728	21	6.76	1.81	6.12	1.88	0.64	1.60	0.35	-0.09	1.37	0.35
FWHM (s)	32	0.906	18	7.89	3.14	6.20	2.40	1.68	2.19	0.52	0.60	2.77	0.60

Data are shown for all patients with PAH initiating or escalating PAH therapy.

DCE, dynamic contrast-enhanced imaging; FWHM, full width at half maximum; ICC, intraclass correlation coefficient; Log₁₀ NT-ProBNP, log to base 10 N-terminal pro-brain natriuretic peptide; 6MWT, six min walk test; PAH, pulmonary arterial hypertension; RVCO, right ventricular cardiac output; RVEDM, right ventricle end-diastolic mass; RVEDV, right ventricle end-systolic volume; RVEF, right ventricle end-systolic volume; RVESM, right ventricle end-systolic mass; RVESV, right ventricle stroke volume; SA, short axis.

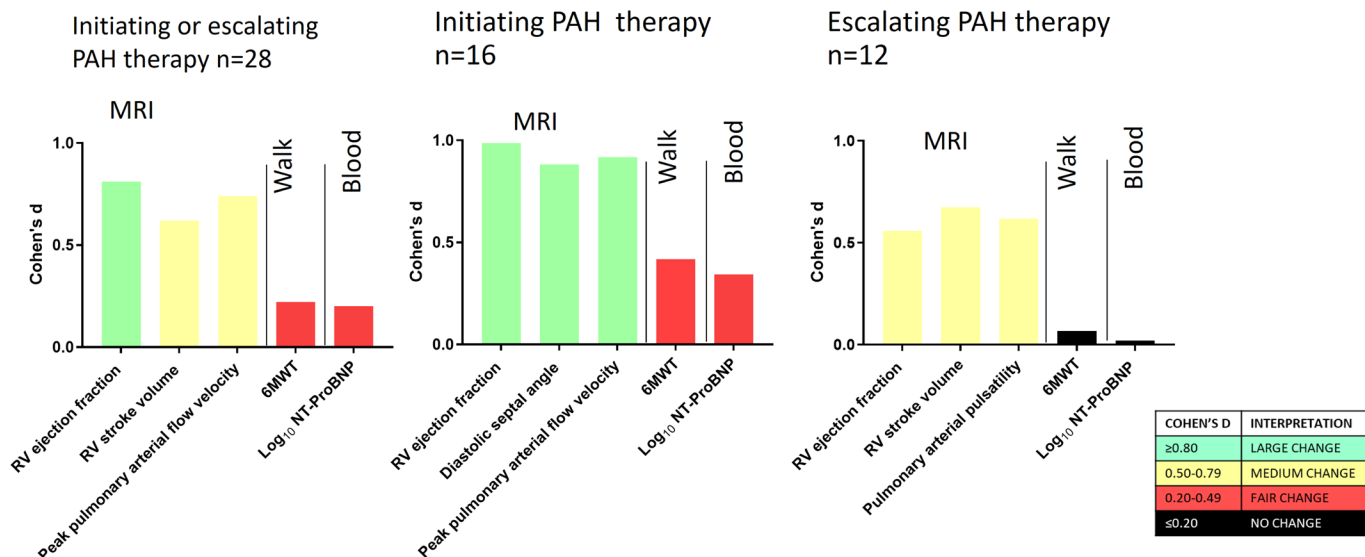


Figure 1 Comparison of treatment effect size using Cohen's d results in patients initiating and/or escalating pulmonary arterial hypertension (PAH) therapy. 6MWT, 6 min walk test; Log₁₀NT-ProBNP, log to base 10 N-terminal pro-brain natriuretic peptide; RV, right ventricular.

Test–test repeatability (visits 2 and 3)

In patients with PAH, test–test repeatability was assessed between visits 2 and 3; 6MWT (ICC 0.987) and log₁₀NT-ProBNP (ICC 0.772) had excellent repeatability. Of cardiac MRI metrics (table 1), all showed excellent repeatability. Data for MRI pulmonary flow and perfusion transit times are shown in table 1.

Treatment effect size (visits 1 and 2)

For all patients, initiating or escalating therapy (n=28), the only measurement with a large treatment effect size was RV ejection fraction (Cohen's d 0.81). The 6MWT (Cohen's d 0.22) and NT-ProBNP (Cohen's d 0.20) demonstrated a fair treatment effect size (table 1). Figure 1 shows Cohen's d values for the

top three MRI end points, the 6MWT and NT-proBNP. Figure 2 shows ICC versus Cohen's d value for all end points. In patients initiating PAH therapy, RV ejection fraction (Cohen's d 0.99), diastolic septal angle (Cohen's d 0.88) and peak pulmonary arterial flow velocity (Cohen's d 0.92) had a large treatment effect size. In patients escalating therapy, RV ejection fraction, RV stroke volume and pulmonary arterial pulsatility had a medium effect size, whereas NT-ProBNP (Cohen's d 0.02) and 6MWT (Cohen's d 0.07) demonstrated no treatment effect (see online supplemental figure S4). The stable patient group showed either no or fair changes across all measured parameters (online supplemental table S6).

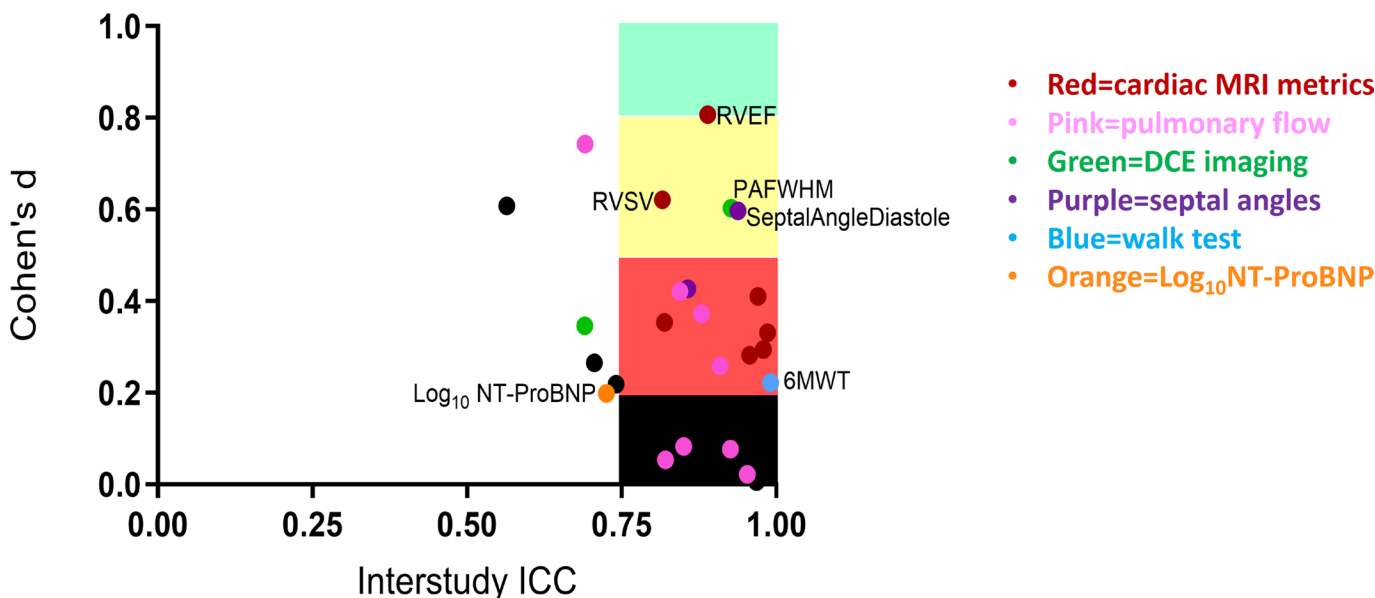


Figure 2 Cohen's d versus interstudy intraclass correlation coefficient (ICC) for study measurements. DCE, dynamic contrast-enhanced imaging; Log₁₀NT-ProBNP, log to base 10 N-terminal pro-brain natriuretic peptide; PAFWHM, pulmonary arterial full width at half maximum; RVEF, right ventricular ejection fraction; RSV, right ventricle stroke volume; 6MWT 6 min walk test. ICC >0.75=excellent repeatability. Cohen's d value of <0.20 was considered no change, 0.20–0.49 was considered fair change, 0.50–0.79 was considered a medium change and ≥0.80 was considered a large change.

DISCUSSION

Investigations used to monitor disease severity in patients with PAH, namely 6MWT distance, NT-ProBNP level and MRI metrics, had excellent repeatability. In contrast, only MRI (RVEF) demonstrated a large treatment effect size in patients initiating or escalating therapies, whereas for the 6MWT and NT-ProBNP the treatment effect sizes were fair.

As observed in previous clinical trials¹ and highlighted at the 6th World Symposium,⁹ all metrics evaluated in patients with PAH escalating therapy had a lower treatment effect size compared with treatment-naïve patients initiating therapy. This represents a challenge when studying the effects of new therapies in PAH where the standard of care is combination treatment.¹ Importantly, MRI was still able to detect a medium treatment effect size in patients receiving background PAH therapy. Due to the large cost of conducting PAH therapy trials, strategies to reduce the size of studies and their duration using a surrogate end point that is repeatable and has a large treatment effect size would be highly desirable.⁹

This study has a number of limitations including the small sample size and the lack of comparison with invasively measured pulmonary haemodynamics. Nonetheless, we have demonstrated in this exploratory study that MRI, the gold standard for RV function assessment, detects a larger treatment effect than the 6MWT or NT-proBNP. This may reflect the ceiling effect of the 6MWT and the effect of comorbidities (including chronic kidney disease) that may influence 6MWT distance and NT-proBNP levels. MRI metrics predict clinical worsening⁷ and mortality^{2–4} fulfilling many of the criteria of a surrogate end point.⁹ Given that pulmonary haemodynamics are commonly used in early phase PAH studies,¹ a direct comparison of MRI metrics and pulmonary haemodynamics, to detect longitudinal change following PAH therapy, is now required if MRI imaging is to be considered a primary end-point for PAH therapy trials.^{8,9}

This study demonstrates the high repeatability of MRI metrics in PAH and the large treatment effect size support further evaluation of MRI as a non-invasive endpoint in PAH therapy trials.

Correction notice This article has been corrected since it was published Online First. Affiliations for FW and AC have been updated.

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Contributors AJS, FW, JMW, AC, LK, DGK conceived the idea for the study. MC, MA, DGK, RC, AJS supported patient recruitment. AJS, JMW, DGK, AM, PH, LS devised the MRI protocol. AJS, CO, PH, LS analysed the MRI studies. FA, AM, CJ, PH, PG performed data quality control checks. MA performed the walk tests and FA, JP, AL performed the lab analyses. MC, LK, SA, AR, PG, AJS, YS, FS, PH, LS supported the data collation and analysis. Statistical analysis was performed by MC, SA, AJS, FA, LS, LK. All authors contributed to the drafting of the manuscript. All authors approved the final version of the manuscript.

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Competing interests FW, LK and AC are employees and shareholders of GlaxoSmithKline. AS is the principal investigator for the collaborative research grant from GlaxoSmithKline that funded this study. AS has undertaken consultancy work for General Electric and Actelion Pharmaceuticals. RC has received fees for lecturing and participation in advisory boards, from Actelion, Bayer, GSK and MSD. DGK has received fees for lecturing and participation in advisory boards, from Actelion, Bayer, GSK and MSD and fees for participation in Steering Committees for Actelion.

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REFERENCES

- Galiè N, Humbert M, Vachiery J-L, *et al*. 2015 ESC/ERS guidelines for the diagnosis and treatment of pulmonary hypertension: the joint Task force for the diagnosis and treatment of pulmonary hypertension of the European Society of cardiology (ESC) and the European respiratory Society (ERS): endorsed by: association for European paediatric and congenital cardiology (AEPCC), International Society for heart and lung transplantation (ISHLT). *Eur Heart J* 2016;37:67–119.
- Swift AJ, Capener D, Johns C, *et al*. Magnetic resonance imaging in the prognostic evaluation of patients with pulmonary arterial hypertension. *Am J Respir Crit Care Med* 2017;196:228–39.
- Lewis RA, Johns CS, Cogliano M, *et al*. Identification of cardiac magnetic resonance imaging thresholds for risk stratification in pulmonary arterial hypertension. *Am J Respir Crit Care Med* 2020;201:458–68.
- van de Veerdonk MC, Kind T, Marcus JT, *et al*. Progressive right ventricular dysfunction in patients with pulmonary arterial hypertension responding to therapy. *J Am Coll Cardiol* 2011;58:2511–9.
- Hurdman J, Condliffe R, Elliot CA, *et al*. ASPIRE registry: assessing the spectrum of pulmonary hypertension identified at a referral centre. *Eur Respir J* 2012;39:945–55.
- Grothues F, Moon JC, Bellenger NG, *et al*. Interstudy reproducibility of right ventricular volumes, function, and mass with cardiovascular magnetic resonance. *Am Heart J* 2004;147:218–23.
- Alabed S, Shahin Y, Garg P, *et al*. Cardiac-MRI predicts clinical worsening and mortality in pulmonary arterial hypertension: a systematic review and meta-analysis. *JACC Cardiovasc Imaging* 2020. doi:10.1016/j.jcmg.2020.08.013. [Epub ahead of print: 30 Sep 2020].
- Kiely DG, Levin D, Hassoun P, *et al*. Express: statement on imaging and pulmonary hypertension from the pulmonary vascular research Institute (PVRI). *Pulm Circ* 2019;2045894019841990. doi:10.1177/2045894019841990
- Sitbon O, Gomberg-Maitland M, Granton J, *et al*. Clinical trial design and new therapies for pulmonary arterial hypertension. *Eur Respir J* 2019;53:1801908.
- Lakens D. Calculating and reporting effect sizes to facilitate cumulative science: a practical primer for t-tests and ANOVAs. *Front Psychol* 2013;4:863.