



Feasibility and reproducibility of a cardiovascular magnetic resonance free-breathing, multi-shot, navigated image acquisition technique for ventricular volume quantification during continuous exercise

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Background: Cardiovascular magnetic resonance (CMR) image acquisition techniques during exercise typically requires either transient cessation of exercise or complex post-processing, potentially compromising clinical utility. We evaluated the feasibility and reproducibility of a navigated image acquisition method for ventricular volumes assessment during continuous physical exercise.

Methods: Ten healthy volunteers underwent supine cycle ergometer (Lode) exercise CMR on two separate occasions using a free-breathing, multi-shot, navigated, balanced steady-state free precession cine pulse sequence. Images were acquired at 3-stages, baseline and during steady-state exercise at 55% and 75% maximal heart rate (HR_{max}), based on a prior supine cardiopulmonary exercise test. Intra- and inter-observer variability and inter-scan reproducibility were derived. Clinical feasibility was tested in a separate cohort of patients with severe mitral regurgitation (n=6).

Results: End-diastolic volume (EDV) of both LV and RV decreased during exercise at 55% and 75% HR_{max} , although a reduction in RVEDV index was only observed at 75% HR_{max} . Ejection fractions (EF) for both ventricles were significantly higher at 75% HR_{max} compared to their respective baselines (LVEF $68\% \pm 3\%$ vs. $58\% \pm 5\%$, $P=0.001$; RVEF $66\% \pm 4\%$ vs. $58\% \pm 7\%$, $P=0.02$). Intra-observer and inter-observer reproducibility of LV parameters was excellent at all 3-stages. Although measurements of RVESV were more variable during exercise, the reproducibility of both RVEF and RV cardiac index was excellent (CV <10%). Inter-scan LV and RV ejection fraction were highly reproducible at all 3 stages, although inter-scan reproducibility of indexed RVESV was only moderate. The protocol was well tolerated by all patients.

Conclusions: Exercise CMR using a free-breathing, multi-shot, navigated cine imaging method allows simultaneous assessment of left and right ventricular volumes during *continuous* exercise. Intra- and inter-observer reproducibility were excellent. Inter-scan LV and RV ejection fraction were also highly reproducible.

Keywords: Cardiovascular magnetic resonance (CMR); exercise; magnetic resonance imaging ergometer (MRI ergometer); free-breathing; feasibility; respiratory-navigation

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1 Introduction

2 Exercise testing can be used to detect underlying
3 cardiovascular abnormalities which are not apparent at
4 rest. Whilst exercise-stress echocardiography and nuclear
5 scintigraphy are widely available, their limitations include
6 poor acoustic windows (1), motion artefacts (2) and
7 radiation exposure (3). Cardiovascular magnetic resonance
8 (CMR) imaging at rest is highly accurate and reproducible
9 (4,5), however exercise stress testing with CMR presents
10 significant challenges. The early evolution of exercise CMR
11 (exCMR) focused on improving the MRI-compatibility of
12 exercise treadmill equipment from being placed external to
13 the MRI room (6), to being in close proximity to the MRI
14 scanner (7-9), to a fully MRI compatible treadmill placed
15 adjacent to the MRI system (8,10-12). These protocols
16 are however limited by the time delay needed to transfer
17 the patient from the treadmill onto the scanner. Any time
18 delay between the cessation of exercise and MRI image
19 acquisition is critical, since exercise-induced functional
20 abnormalities may begin to disappear almost immediately
21 after exercise cessation (13-15).

23 The development of a MRI-compatible cycle ergometer
24 allows patients to exercise in the supine position whilst
25 inside the bore of the magnet (16). Imaging during
26 continuous exercise eliminates the time lapse between
27 exercise and imaging and may allow a more accurate
28 assessment of changes in cardiac physiology during exertion.
29 Excessive motion during exercise however poses a challenge
30 in image acquisition. As a result, investigators have
31 resorted to acquire images following transient cessation of
32 exercise (17), during breath-holds (6,17,18) or using
33 ungated real-time cine imaging (19). Reconstruction of a
34 short axis stack for volumetric analysis from ungated real-
35 time imaging, however, involves complex post-processing
36 analysis in addition to a requirement for bespoke in-house
37 software (19).

38 The objectives of this study were: (I) to assess the
39 feasibility and reproducibility of a navigated cine image
40 acquisition method for the assessment of the ventricular
41 volumes during continuous exercise; (II) to examine its
42 clinical feasibility in patients with significant valvular heart
43 disease.

45 Methods

47 Study design and population

48 This study was performed in 3 stages: (I) a pilot phase

in which the feasibility of a navigated image acquisition
sequence was tested in healthy volunteers; (II) an assessment
of inter-scan reproducibility in which each healthy volunteer
underwent a repeat exCMR after a median of 16 weeks;
(III) clinical application of this technique in patients with
severe mitral regurgitation (MRegur). The study was
approved by a local ethics committee (Yorkshire & The
Humber-Leeds West 12/YH/0551) and complied with the
Declaration of Helsinki. All participants provided written
informed consent.

Pilot phase & reproducibility

Ten healthy volunteers with no history or symptoms of
cardiovascular disease and no contraindications to CMR
were recruited. Absolute and relative contraindications to
exercise testing were adhered to according to American
Heart Association (AHA) guidelines (20). All participants
had a height of <190 cm. All healthy volunteers underwent
a supine cardiopulmonary exercise test (CPET) prior to
undertaking exCMR on a supine cycle ergometer. CMR
was performed on a 1.5 Tesla MRI system with 70 cm bore
(Ingenia, Philips Healthcare, Best, Netherlands) equipped
with a 28-channel coil and free-breathing images were
acquired during continuous exercise. Exercise intensity
was individualized to the heart rate (HR) corresponding to
55% and 75% of the maximal HR (HR_{max}) attained on their
pre-CMR supine CPET. After a median time of 16 weeks,
exCMR was repeated using an identical scanner and
protocol.

Clinical feasibility

The potential for translation of this technique into clinical
practice was examined in a separate cohort of 6 patients
with significant MRegur, all prospectively recruited from
the valvular heart disease clinic at Leeds Teaching Hospitals
NHS Trust. Inclusion criteria included: moderate-severe or
severe MRegur on echocardiography, and New York Heart
Association functional Class I. Exclusion criteria included:
contraindications to exercise stress testing according to
AHA guidelines (20), presence of atrial fibrillation, height
>190 cm, inability to exercise and contraindications to
CMR. In our institution treadmill CPET is used clinically
in patients with significant MRegur and we utilized these
data to prescribe the individualized HR during exCMR.
To allow for the lower HR response in supine cycling
compared to upright treadmill exercise and the reduced

98 exercise tolerance seen in patients with severe MRegur, the
 99 prescribed HR had to be altered from healthy volunteers.
 100 Patients were thus exercised to 30–39% and 40–59% of
 101 their heart rate reserve (HRR), corresponding to ‘light’ and
 102 ‘moderate’-intensity exercise according to the American
 103 College of Sports Medicine guidelines (21). HRR was
 104 calculated based on this formula: resting HR on CPET +
 105 [$x\%$ of (max HR achieved on treadmill CPET – resting
 106 HR)]; where x is the target % of HRR.

107

108

109 *Cardiopulmonary exercise testing*

110

111 All healthy volunteers underwent CPET on a supine cycle
 112 ergometer (Lode BV, Groningen, The Netherlands).
 113 The crank length on the pre-CMR cycle ergometer was
 114 adjusted to replicate the setup of the in-scanner MRI
 115 ergometer. CPET was conducted as a ramp incremental test
 116 (15 W/min) to volitional intolerance. Breath-by-breath
 117 analysis of the volume and concentration of expired gases
 118 was achieved using an automated system (Medgraphics
 119 Ultima, Minnesota, USA). HR was continuously monitored
 120 via an attached 12-lead electrocardiogram (ECG). The main
 121 outcome measures were maximal HR and maximal power
 122 output in Watts. ExCMR was performed after a median of
 123 8 days [interquartile range (IQR) 2–13].

123

124

125 *Exercise CMR protocol and image acquisition*

126

127 Exercise whilst in the bore of the magnet was conducted
 128 on a supine MRI-compatible cycle ergometer (Lode
 129 BV, Groningen, The Netherlands). Optimal participant
 130 preparation included instructions on consistent thoracic
 131 breathing, use of handrail to ensure trunk stability, skin
 132 preparation to maximize interface between electrode and
 133 skin, and securing vector ECG connections onto anterior
 134 chest wall with tape to ensure quality recording of ECG. A
 135 blood pressure (BP) cuff was placed on the left arm. Both
 136 the surface coil and torso pad were then firmly secured onto
 137 the participants with elastic Velcro® straps. The MRI table
 138 was advanced whilst participants performed a short bout of
 139 unloaded exercise to ensure that their knees did not contact
 the scanner casing during pedalling.

140 Free-breathing images were acquired at 3-stages, at
 141 rest and then during steady-state exercise at 55% HR_{max}
 142 and 75% HR_{max} . Exercise began with a 2 min warm-
 143 up at a power output of 0 W (unloaded). Work rate was
 144 incrementally increased by 10–20 W until the target 55%
 145 HR_{max} was achieved, and then adjusted to maintain the

HR at the required target throughout the exercise. Verbal
 feedback was constantly given to participants and cycling
 cadence was maintained between 60–70 rpm. Following
 a rest period of 2 minutes, a second bout of exercise was
 undertaken until the target 75% HR_{max} was achieved.
 Heart rate and rhythm were continuously monitored, and
 BP was recorded at each stage. Each stage of exercise was
 maintained for 5–7 minutes (2 minutes to achieve steady-
 state in HR and approximately 3–5 minutes of image
 acquisition). Imaging was only performed during steady-
 state conditions, when HR was maintained at near constant
 levels. Criteria for termination prior to achieving target
 HR included participant’s request and a drop in systolic BP
 >10 mmHg.

The scan protocol included standard long axis views
 (vertical, horizontal long axis) and a short axis ventricular
 volume stack. Cine imaging was performed using a free-
 breathing, multi-shot, respiratory-navigated, balanced
 steady-state free precession pulse sequence. A respiratory
 echo-based navigator was placed on the right hemi-
 diaphragm with a 5 mm gating window and continuous
 gating level drift activated. A cylindrical MR radiofrequency
 excitation pulse from which a 1-dimensional projection of
 the lung-liver interface was generated and was used to infer
 the breathing phase. The navigator was played at the start
 of the R-R interval, at end-diastole of the cardiac cycle.
 The steady-state of ongoing balanced steady-state free
 precession (bSSFP) readout was stopped in the standard
 controlled manner by using half-alpha radiofrequency
 pulses to temporarily store the steady state magnetization
 in the z-direction. This allowed the respiratory navigator
 to last for a total duration of 24 ms (played out for 17 ms
 before resuming readout after 7 ms), equivalent to
 approximately 9 repetition time (TR). Retrospective
 cardiac triggering was used in this study (continuous data
 sampling). The bSSFP readout was continuous, wherein
 data from the ECG and k-space profile acquisition timings
 were matched to produce images for all cardiac phases.
 Cartesian sampling was used, and the acquired k-space lines
 were only accepted for image reconstruction if the right
 hemi-diaphragm position was within the gating window
 during end-expiratory phase. K-space profiles which were
 rejected outside of the respiratory navigator gate were
 reacquired. Other scan parameters were as follows: typical
 field of view (FOV) 320 mm × 320 mm, repetition time (TR)
 2.8 ms, echo time (TE) 1.4 msec, flip angle 60°, temporal
 resolution 33 ms, SENSE factor 2, multi-shot turbo field
 echo (TFE) factor 11, TFE acquisition duration 30.4 ms,

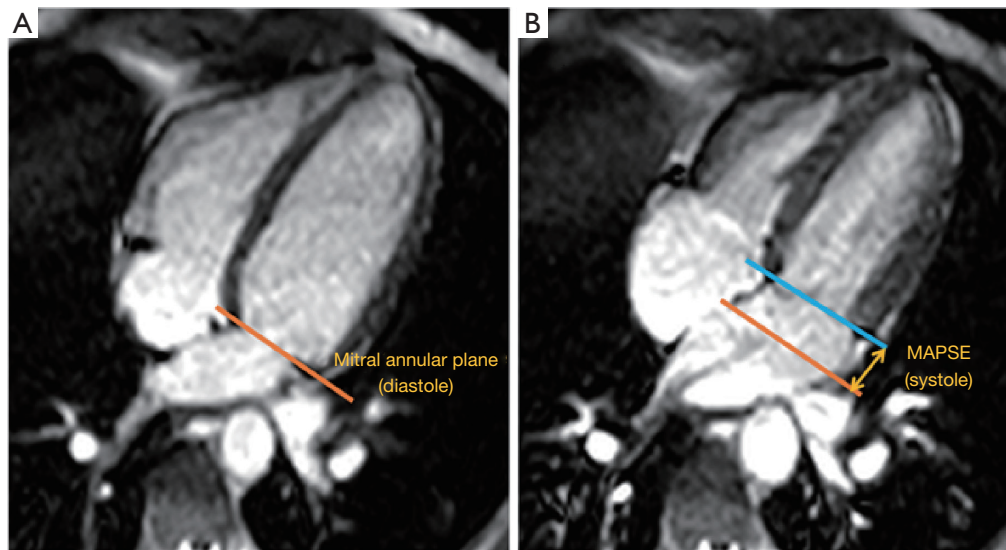


Figure 1 Assessment of longitudinal function. (A) End-diastole phase was identified and a reference line (orange) was drawn across the atrioventricular valve plane and forwarded across all phases of the cine image. (B) A further line (blue) is drawn in end-systole. The distance between the two points at the lateral mitral valve annulus (MAPSE) was measured and expressed in mm. MAPSE, mitral annular plane systolic excursion.

194 phase percentage 50%, slice thickness 10 mm, 0 mm gap,
 195 in-plane spatial resolution 2.4 mm × 2.4 mm and matrix
 196 132×106. A total of 16 cardiac phases were acquired and this
 197 was reconstructed to 30 cardiac phases.

198

199 *CMR analysis*

200

201 CMR analysis was performed by two independent operators
 202 (PC, LB; both observers with 3 years CMR experience)
 203 using commercially available computer software (cmr⁴²,
 204 Circle Cardiovascular Imaging Inc, Calgary, Alberta,
 205 Canada). Left and right ventricular volumes, and ejection
 206 fraction (EF) were calculated in the conventional method,
 207 by manually tracing endocardial contours in end-diastole
 208 and end-systole on the short axis stack (*Figure S1*).
 209 Biventricular end-diastolic and end-systolic volumes were
 210 calculated using a summation of discs technique (22). Stroke
 211 volume (SV) was measured as the difference between end-
 212 diastolic and end-systolic volume, whereas cardiac output
 213 was calculated as: $SV \times HR$. All measured volumes and
 214 cardiac output parameters were indexed to body surface area
 215 (Mostellar formula). Longitudinal LV function in the form
 216 of mitral annular plane systolic excursion (MAPSE) was
 217 assessed by using mitral annular excursion. In the 4-chamber
 218 cine image, atrioventricular motion was measured at the

lateral junction points between the left atrium and ventricle
 at end diastole and end systole. The perpendicular distance
 between these two points was measured. *Figure 1* outlines
 the methodology used to assess LV longitudinal contraction.

Statistical analysis

All statistical analysis was performed using the SPSS V.21.0
 (IBM Corp., New York, USA). All continuous data were
 tested for normality using the Shapiro-Wilk test; variables
 are expressed as mean ± SD or median (IQR) in cases of
 skewed distributions. Categorical variables are expressed as
 frequencies and percentages. Repeated measures analysis
 of variance (ANOVA) with Bonferroni post-test analysis
 was used to compare data between rest and different stages
 of exercise. Intra- and inter-observer reproducibility was
 assessed by the coefficient of variation (CV) test, the
 standard deviation of differences between observations
 divided by the mean. $P < 0.05$ was considered statistically
 significant.

Results

Healthy volunteers and baseline CMR data

All 10 healthy volunteers [7 men, age 25 ± 2 years, body

Table 1 Volumetric data at baseline, and during exercise at 55% and 75% HR_{max} in healthy volunteers

Cardiovascular variables	Baseline	55% HR _{max}	75% HR _{max}	P value (baseline vs. 55% HR _{max})	P value (55% HR _{max} vs. 75% HR _{max})	P value (baseline vs. 75% HR _{max})
LVEDV (mL)	182±28	175±27	159±22	0.003	0.010	0.001
LVEDV (indexed), mL/m ²	97±11	93±10	85±7	0.002	0.012	0.001
LVESV (mL)	77±18	68±19	52±9	0.269	0.022	0.001
LVESV (indexed), mL/m ²	41±7	36±9	28±3	0.252	0.019	0.001
LVSV (mL)	105±14	107±21	107±15	1.000	1.000	1.000
LVSV (indexed), mL/m ²	57±6	57±10	57±5	1.000	1.000	1.000
LVEF (%)	58±5	61±8	68±3	0.912	0.109	0.001
LV cardiac output, mL/min	7,087±1,392	10,188±2,902	14,041±2,454	0.004	0.005	<0.001
LV cardiac index, mL/min/m ²	3,805±721	5,456±1,448	7,503±1,055	0.003	0.003	<0.001
RVEDV (mL)	178±30	171±182	152±25	0.257	0.022	0.011
RVEDV (indexed), mL/m ²	95±11	92±8	81±7	0.231	0.020	0.009
RVESV (mL)	76±21	66±18	52±12	0.119	0.134	0.017
RVESV (indexed), mL/m ²	40±10	35±8	28±5	0.124	0.129	0.011
RVSV (mL)	102±17	105±14	101±16	1.000	1.000	1.000
RVSV (indexed), mL/m ²	51±9	56±5	54±5	0.270	1.000	0.872
RVEF (%)	58±7	62±7	66±4	0.365	0.463	0.017
RV cardiac output, mL/min	6,869±1,752	9,957±2,327	13,119±2,196	0.002	0.009	<0.001
RV cardiac index, mL/min/m ²	3,685±907	5,333±1,133	6,991±704	0.002	0.007	<0.001

Data as mean ± SD. LVEDV, left ventricular end-diastolic volume; LVESV, left ventricular end-systolic volume; LVSV, left ventricular stroke volume; LVEF, left ventricular ejection fraction; LV, left ventricle; RVEDV, right ventricular end-diastolic volume; RVESV, right ventricular end-systolic volume; RVSV, right ventricular stroke volume; RVEF, right ventricular ejection fraction; RV, right ventricle.

245 mass index (BMI) 23.1±2.2 kg/m² completed the full study
 246 protocol. HR increased during exercise (68±12 vs. 94±13
 247 vs. 131±11 bpm, baseline vs. 55% HR_{max} vs. 75% HR_{max};
 248 all P<0.001). Systolic BP was significantly higher during
 249 exercise at 75% HR_{max} than at baseline (130±12 vs. 120±
 250 10 mmHg; P=0.03), whilst diastolic BP remained unchanged
 251 (70±14 vs. 70±8 mmHg; P=1.00). Mean supine work rate for
 252 exercise at 55% HR_{max} and 75% HR_{max} was 25±19 W and
 253 87±23 W, respectively. CMR data for all subjects are shown
 254 in *Table 1*. *Figure 2* demonstrated the exCMR images at
 255 baseline and during exercise.

256

257

Left ventricular (LV) and right ventricular (RV)

parameters during exercise

258

259

260 The changes in ventricular volumes during exercise are

plotted in *Figure 3*. End-diastolic volume (EDV) of the LV 261
 decreased significantly during exercise at 55% and 75% 262
 HR_{max}. In contrast, RVEDV remained unchanged from 263
 baseline at 55% HR_{max} and significantly decreased at 75% 264
 HR_{max} (P=0.02). LV end-systolic volume (LVESV) decreased 265
 when exercised from 55% HR_{max} to 75% HR_{max} (P=0.02). 266
 During exercise at 55% HR_{max}, LVESV was however 267
 not significantly different from baseline. RV end-systolic 268
 volume (RVESV) significantly decreased during exercise at 269
 75% HR_{max} compared to baseline. Both LV and RV stroke 270
 volumes remained unchanged. Ejection fractions (EF) for 271
 both ventricles were significantly higher during exercise 272
 at 75% HR_{max} when compared to their respective baseline 273
 values (LVEF 68%±3% vs. 58%±5%; P=0.001 and RVEF 274
 66%±4% vs. 58%±7%; P=0.02). During exercise, LV and 275
 RV cardiac indexes also increased significantly (*Figure 4*). 276

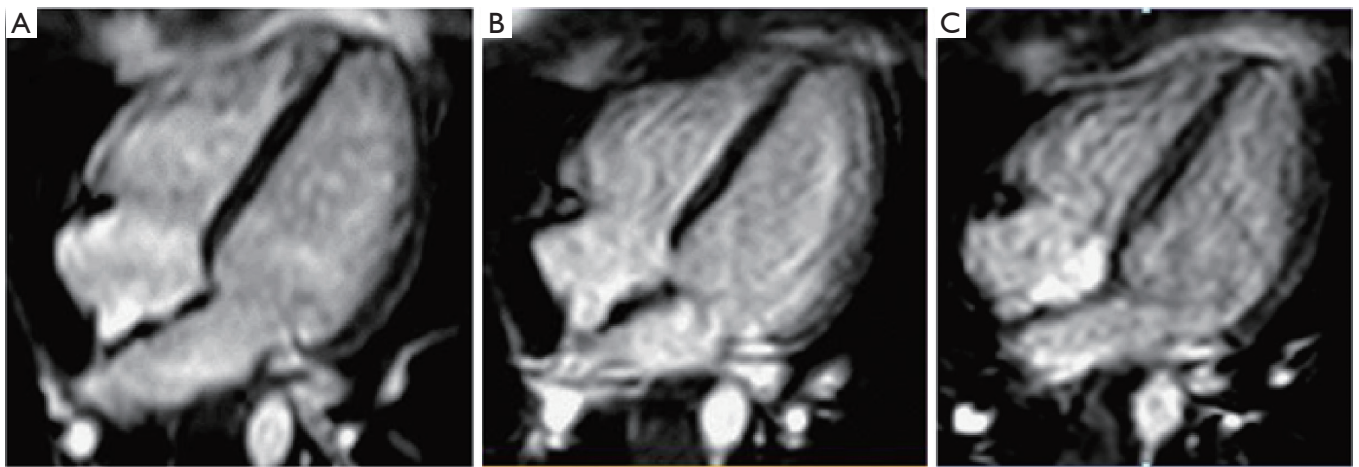


Figure 2 exCMR images at baseline (A), and during exercise at 55% HR_{max} (B) and 75% HR_{max} (C). HR_{max} maximal heart rate.

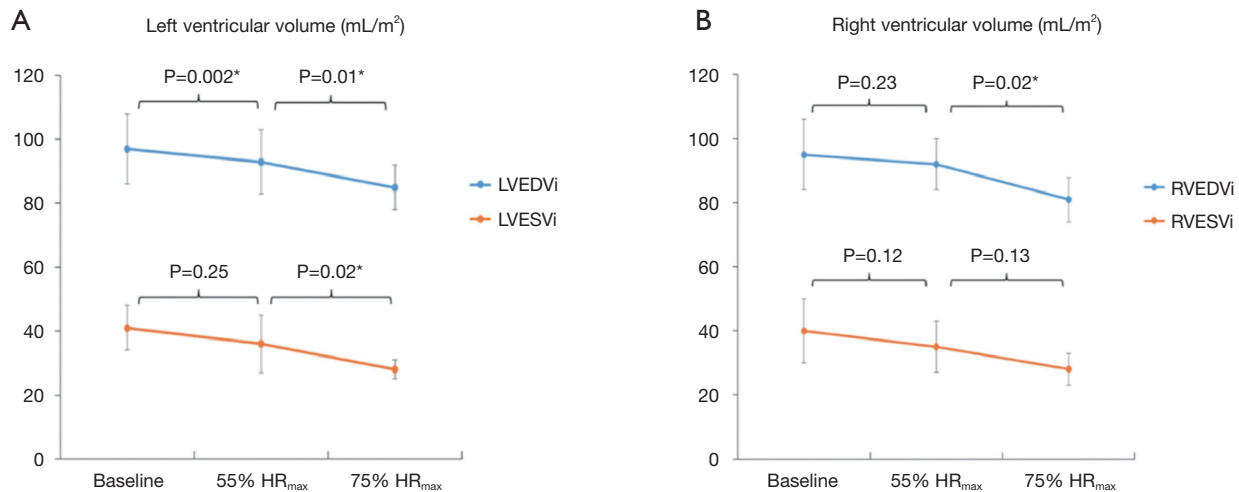


Figure 3 Ventricular volumes during exercise in healthy volunteers. LV (A) and RV (B) end-diastolic and end-systolic volumes during exercise in healthy volunteers. LVEDVi, indexed LV end-diastolic volume; LVESVi, indexed LV end-systolic volume; RVEDVi, indexed right ventricular end-diastolic volume; RVESVi, indexed right ventricular end-systolic volume; HR_{max} , maximal heart rate.

277 *LV longitudinal contraction*

278 Baseline MAPSE appears to be higher in healthy volunteers
 279 when compared to patients with severe mitral regurgitation
 280 (14 ± 4 vs. 12 ± 3 mm) (Figure 5). In the healthy volunteers,
 281 MAPSE increased from 14 ± 4 to 19 ± 5 mm ($P=0.05$) during
 282 exercise at 55% HR_{max} . At 75% HR_{max} , MAPSE appears
 283 to decrease to 17 ± 4 mm although this change was not
 284 statistically significant ($P=1.00$). In patients with severe
 285 mitral regurgitation, there was no significant change of
 286 MAPSE between baseline and both stages of exercise.
 287

288 There is a trend however, indicating that MAPSE increased
 289 with exercise and appear to decline slightly when higher
 290 intensity exercise was achieved.
 291

292 *Intra- and inter-observer reproducibility*

293
 294 Intra-observer reproducibility of LV volumes, LV ejection
 295 fraction and LV cardiac index was excellent at all three
 296 stages, evidenced by $CV \leq 10\%$ (Table 2). During exercise,
 297 the measurements of RVESV were more variable (CV
 298 11–20%). The reproducibility of RV EDV, RV ejection

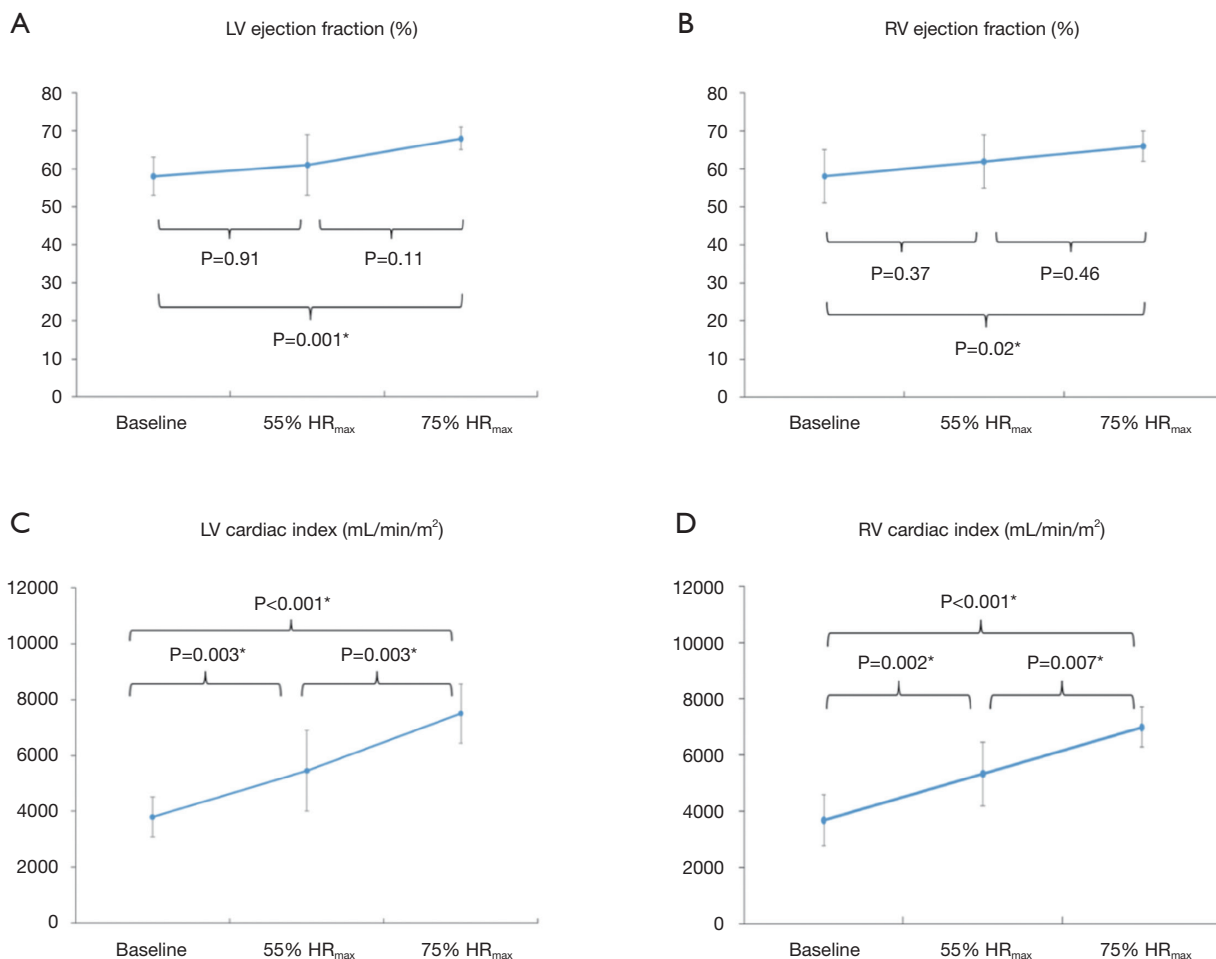


Figure 4 Exercise cardiac reserve in healthy volunteers. (A,B) LV and RV ejection fraction; (C,D) LV and RV cardiac indexes during exercise in healthy volunteers. Data presented in mean (dots) and standard deviation (bars). Asterisks denote statistically significant differences ($P < 0.05$). HR_{max}, maximal heart rate.

Group	MAPSE (baseline)	MAPSE (55% HR _{max})	MAPSE (75% HR _{max})	P value (baseline vs. 55% HR _{max})	P value (55% HR _{max} vs. 75% HR _{max})	P value (baseline vs. 75% HR _{max})	ANOVA with Bonferroni correction
Healthy volunteers (n=10)	14±4 mm	19±5 mm	17±4 mm	0.05	1.00	0.24	0.04
Group	MAPSE (baseline)	MAPSE (light intensity)	MAPSE (moderate intensity)	P value (baseline vs. light)	P value (light vs. moderate)	P value (baseline vs. moderate)	ANOVA with Bonferroni correction
Patients with MR (n=5)	12±3 mm	17±4 mm	15±4 mm	0.27	1.00	0.76	0.21

299 **Figure 5** Longitudinal contraction in both healthy volunteers and patients with severe mitral regurgitation. Data as mean ± SD. MAPSE, 304
 300 mitral annular plane systolic excursion; MR, mitral regurgitation. 305
 301 306
 302 307
 303 308

309 fraction, and RV cardiac index was however excellent (CV
310 <10%).

311 Inter-observer reproducibility of LV volumes, LV
312 ejection fraction and LV cardiac index was also excellent
313 at all three stages (CV for LVEDV $\leq 5\%$; LVESV $\leq 10\%$;
314 LVEF <6%; LV cardiac index <8%). With incremental
315 exercise, inter-observer reproducibility was better in
316 the assessment of RVEDV (CV <5%), when compared
317 to RVESV measurements (CV 12–14%). Although
318 measurements of RVESV were more variable during
319 exercise, the reproducibility of RV ejection fraction, RV
320 stroke volume and RV cardiac index was however excellent.
321 During exercise at 75% HR_{max}, inter-observer LVESV was
322 more reproducible than RVESV (CV 10% vs. 14%).
323

324 *Inter-scan reproducibility*

325
326 We observed good inter-scan reproducibility for LV end-
327 diastolic and end-systolic volumes during exercise; although
328 only modest reproducibility was seen in the readings of LV
329 cardiac index (CV 10–16%). The RVESV measurements
330 were the least reproducible (CV 11–24%). Inter-scan LV
331 and RV ejection fraction were however highly reproducible
332 (CV <10%) at all 3 stages.
333

334 *Ventricular volumes in clinical patients*

335
336 Of 6 patients with severe MRegur, 5 patients (60% men, age
337 60±14 years, BMI 24±2.2 kg/m²) completed the full study
338 protocol. exCMR had to be abandoned in 1 patient due to a
339 significant hypotensive response. HR increased throughout
340 exercise (73±6 vs. 111±11 vs. 118±18 bpm, baseline vs. light
341 vs. moderate; all P<0.01). Systolic BP was significantly
342 higher during moderate intensity exercise than at baseline
343 (114±6 vs. 148±15 mmHg; P=0.02), whilst diastolic BP
344 remained constant (74±9 vs. 80±8 vs. 66±14 mmHg;
345 P=1.00). Mean supine work rate for light and moderate-
346 intensity exercise was 44±19 and 53±32 W, respectively.
347 CMR data for all clinical patients are described in *Table 3*.
348 There was no significant change in the LVEDV during
349 exercise in this small patient sample, and despite a
350 downward trend of LVESV, this was not significant. LVEF
351 was significantly higher when moderate-intensity exercise
352 was achieved. The augmentation of cardiac output and
353 cardiac index was apparent with incremental exercise.
354 When considering the RV parameters, there was no
355 significant change in its EDV. During moderate-intensity
exercise, RVESV was significantly smaller than at baseline.

356 Despite a numerical increase in RVEF with exercise, this
357 was not significant. RV cardiac output and RV cardiac index
358 were unchanged during light-intensity exercise but were
359 significantly increased during moderate-intensity exercise.
360

361 **Discussion**

362
363 This study demonstrated the (I) feasibility of the free-
364 breathing, multi-shot, navigated image acquisition method
365 in the serial assessment of ventricular volumes during
366 continuous exercise; (II) excellent intra- and inter-observer
367 reproducibility, in particular the LV indices; (III) clinical
368 feasibility of this imaging method in a challenging group
369 of patients with significant mitral regurgitation, the first
370 exCMR study performed in this patient group.

371 Previously, image acquisition techniques using the MRI
372 cycle ergometer have either involved a brief period of
373 exercise cessation (17) or required a breath-hold protocol
374 (6,18) in order to reduce excessive motion artefacts and
375 avoid poor ECG signal. Ungated real-time CMR imaging
376 (19,23,24) has been a method that enabled cine images to
377 be acquired during continuous exercise. However, the post-
378 processing analysis of these images requires retrospective
379 synchronization of ECG and respiratory movements, in
380 addition to the need for non-commercially available in-
381 house software (19), therefore decreasing widespread
382 attainability. The application of other image acquisition
383 techniques such as motion correction (25,26) can be
384 challenging in this setting due to the large amount of
385 through plane motion during exercise. Navigator-echo-
386 based gating techniques have been practical methods
387 for effective reduction of respiration motion effects, and
388 are well established for coronary MRI imaging (27,28).
389 Our feasibility study demonstrated that the application
390 of respiratory-navigated technique in exCMR has the
391 potential to overcome respiratory motion which can be
392 quite significant during vigorous exercise. This technique
393 was feasible in both healthy volunteers and clinical patients,
394 and the images acquired were analyzable and reproducible.
395 Moreover, this imaging technique allowed serial assessment
396 of cardiac function at incremental exercise with a further
397 advantage that image analysis can be performed on widely
398 used, commercially available software. This protocol
399 therefore has the potential to increase the utility of exCMR
400 as a clinical assessment tool.

401 La Gerche *et al.* (19) compared real-time ungated with
402 gated CMR techniques and demonstrated that despite
its complex post-processing analysis, ventricular volumes

Table 2 Coefficient of variation (CV) for the reproducibility of LV and RV cardiac indices

Stages	Cardiovascular variables	Coefficient of variation for reproducibility (%)		
		Intra-observer	Inter-observer	Inter-scan
REST	LVEDVi	3.3	2.6	7.6
	LVESVi	8.1	7.3	6.8
	LVSVi	4.3	6.4	12.7
	LVEF	4.5	5.4	6.5
	LV CI	4.3	5.3	15.1
	RVEDVi	4.3	4.8	7.1
	RVESVi	9.6	9.8	15.1
	RVSVi	8.5	6.5	11.4
	RVEF	6.8	5.1	10.3
	RV CI	8.5	5.7	17.2
Exercise at 55% HR _{max}	LVEDVi	3.2	2.7	5.5
	LVESVi	10.0	6.5	11.7
	LVSVi	5.1	5.3	12.5
	LVEF	5.6	3.7	9.2
	LV CI	5.1	5.3	16
	RVEDVi	5.5	4.6	8.3
	RVESVi	11.6	12.4	16.1
	RVSVi	6.3	5.8	9.5
	RVEF	5.1	6.0	7.1
	RV CI	6.3	6.0	12.3
Exercise at 75% HR _{max}	LVEDVi	6.4	4.8	7.1
	LVESVi	9.8	10	11.6
	LVSVi	9.3	7.3	10.1
	LVEF	4.9	5.3	5.8
	LV CI	9.1	7.1	10.1
	RVEDVi	6.6	3.5	12.1
	RVESVi	19.5	13.6	23.5
	RVSVi	8.4	4.9	10.4
	RVEF	7.7	5.5	8.5
	RV CI	8.5	4.8	8.8

Data as %. HR_{max}, maximal heart rate; LVEDVi, indexed left ventricular end-diastolic volume; LVESVi, indexed left ventricular end-systolic volume; LVSVi, indexed left ventricular stroke volume; LVEF, left ventricular ejection fraction; LV CI, left ventricular cardiac index; RVEDVi, indexed right ventricular end-diastolic volume; RVESVi, indexed right ventricular end-systolic volume; RVSVi, indexed right ventricular stroke volume; RVEF, right ventricular ejection fraction; RV CI, right ventricular cardiac index; HR, heart rate.

Table 3 Volumetric data at baseline, and during light and moderate-intensity exercise in clinical patients

Cardiovascular variables	Baseline	Light intensity	Moderate intensity	P value (baseline vs. light)	P value (light vs. moderate)	P value (baseline vs. moderate)
LVEDV (mL)	187±42	187±41	184±48	1.00	1.00	1.00
LVEDV (indexed), mL/m ²	96±25	95±23	94±26	1.00	1.00	1.00
LVESV (mL)	78±19	64±15	62±20	0.24	0.24	0.12
LVESV (indexed), mL/m ²	40±11	33±8	32±11	0.20	0.21	0.14
LVSV (mL)	109±27	123±29	122±29	0.07	0.07	0.10
LVSV (indexed), mL/m ²	56±15	63±16	62±16	0.09	0.09	0.11
LVEF (%)	58±4	65±5	67±3	0.08	0.08	0.04
LV cardiac output, mL/min	8,081±2,570	13,723±3,719	14,460±3,957	0.003	0.003	0.004
LV cardiac index, mL/min/m ²	4,129±1,389	7,022±2,107	7,406±2,296	0.01	0.01	0.01
RVEDV (mL)	181±60	176±64	176±62	1.00	1.00	1.00
RVEDV (indexed), mL/m ²	91±30	89±32	89±30	1.00	1.00	1.00
RVESV (mL)	85±30	85±41	67±24	1.00	1.00	0.02
RVESV (indexed), mL/m ²	43±15	43±21	34±12	1.00	1.00	0.03
RVSV (mL)	96±30	91±29	109±40	1.00	1.00	0.91
RVSV (indexed), mL/m ²	48±15	46±14	55±19	1.00	1.00	1.00
RVEF (%)	53±4	53±10	62±5	1.00	1.00	0.13
RV cardiac output, mL/min	7,112±2,865	10,120±3,374	12,544±3,933	0.15	0.15	0.01
RV cardiac index, mL/min/m ²	3,609±1,449	5,136±1,750	6,355±1,952	0.15	0.15	0.01

Data as mean ± SD. LVEDV, left ventricular end-diastolic volume; LVESV, left ventricular end-systolic volume; LVSV, left ventricular stroke volume; LVEF, left ventricular ejection fraction; LV, left ventricle; RVEDV, right ventricular end-diastolic volume; RVESV, right ventricular end-systolic volume; RVSV, right ventricular stroke volume; RVEF, right ventricular ejection fraction; RV, right ventricle.

403 were analysable more frequently with real-time ungated
 404 compared with gated CMR (100% vs. 47%; P<0.001). In
 405 our gated CMR study, when combined with ‘respiratory-
 406 navigation’, sufficient image quality for analysis was achieved
 407 in 100% of the scans. La Gerche *et al.* also observed better
 408 interobserver variability for real-time ungated (CV =1.9%
 409 and 2.0% for LV and RV stroke volumes, respectively)
 410 than gated scans (CV =15.2% and 13.6%; P<0.01) (19).
 411 Comparing their gated study to ours, the incorporation of
 412 ‘respiratory-navigation’ in our gated study improved the
 413 CV for left and RV stroke volumes (CV of 7.3% and 4.9%,
 414 during exercise at 75% HR_{max}, respectively).

415 In 2017, Le *et al.* (24) combined real-time imaging with
 416 ECG-gated sequences to assess exercise cardiac volumetrics
 417 in healthy volunteers and athletes. Image acquisition,
 418 however, required suspension of exercise at the end of

every stage for free-breathing imaging. The decline in 419
 HR following cessation of exercise (15,29) can potentially 420
 impair diagnostic accuracy and clinical utility. In contrast, 421
 our exCMR protocol permits imaging during continuous 422
 exercise, eliminating the time lapse between exercise and 423
 imaging altogether. In relation to scan parameters, our 424
 study had better temporal resolution (33 vs. 39 ms) and a 425
 smaller voxel size (2.4 mm × 2.4 mm vs. 3.3 mm × 2.3 mm) 426
 indicating improved spatial resolution. 427

The effects of left and RV volumes during physical 428
 exercise remains controversial. Some data are generally 429
 consistent with an enhanced contractile state during supine 430
 exercise, but the role of Frank Starling mechanism remains 431
 uncertain. In supine exCMR, there are several factors that 432
 affect the hemodynamic response. Firstly, exercise in the 433
 supine position results in a lower HR response but a greater 434

435 rise in systolic BP, compared with upright exercise (30,31).
436 This results in a similar double product ($HR \times$ systolic
437 BP), which is an index of myocardial oxygenation (32),
438 and a rationale for similar detection rates of ischemia
439 between upright and supine stress echocardiography despite
440 a significant lower peak heart rates with supine exercise
441 (30,33). As such, for a given exercise intensity, lower heart
442 rates are expected in supine exercise, compared with upright
443 exercise. Secondly, due to increased afterload in the supine
444 position, it is postulated that end-systolic volumes are near
445 maximal even at rest. As a result, a recent meta-analysis
446 in exCMR studies by Beaudry *et al.* demonstrated no
447 significant rise in LVEDV, with many studies demonstrating
448 a non-significant decrease with exercise (34). This meta-
449 analysis also demonstrated an exercise rise in LV stroke
450 volume driven by a fall in end-systolic volume. The authors
451 however did not account for one important factor, which is
452 the effect of respiration on cardiac hemodynamics. Claessen
453 *et al.* (35) elegantly demonstrated the significant effect
454 respiration cycles have on cardiac hemodynamics; with end-
455 expiration resulting in a significant rise in LVEDV and LV
456 stroke volume and a fall in RVEDV and RV stroke volume,
457 when compared with end-inspiration at numerous levels of
458 exercise. The results of this meta-analysis should therefore
459 be interpreted with caution, as although insightful, it
460 analyses a heterogeneous group of exCMR studies including
461 both free-breathing and breath-held acquisitions, in a varied
462 mix of healthy volunteers, endurance athletes and patients
463 with cardiac disease, and who were exercised to different
464 exercise intensities.

465 The results of this present study are in line with previous
466 studies of supine exercise, showing a decrease in LV
467 (24,36) and RV (18,23,37) EDVs, particularly during later
468 stages of exercise. Similar to previous exCMR studies, we
469 demonstrated no significant rise in stroke volume with
470 exercise (38). Healthy volunteers have been shown to achieve
471 their peak diastolic filling and contractility earlier (24).
472 As a result, LVEDV in healthy volunteers peaked earlier
473 and decreased subsequently. The increase in HR during
474 exercise also reduced diastolic filling time, therefore leading
475 to smaller LV and RV cavity during diastole. It is worth
476 noting that as this study assessed 2 stages of exercise (55%
477 and 75% HR_{max}) at moderate and high exercise intensities
478 respectively, it is possible our data did not capture the initial
479 LV dilatation described in the Frank Starling mechanism.
480 Indeed, numerous prior investigators (23,24,35) have
481 demonstrated an initial rise in LVEDV and LV stroke

482 volume at early lower exercise stages followed by a fall at
483 later higher intensity stages. However, due to the nature
484 of exCMR studies being performed in small numbers,
485 these changes often do not reach statistical significance. As
486 such it is likely our study did not capture this early rise in
487 LVEDV and LV stroke volume given the exercise stages
488 were performed at moderate and high intensities. This is
489 the first study to confirm clinical feasibility of this exCMR
490 protocol in patients with severe MRegur. Ventricular
491 volumes in patients were unchanged during light and
492 moderate intensity exercise, which is likely a reflection of
493 their relatively deconditioned state and poorer response in
494 terms of myocardial contractility.

495 CMR MAPSE has been proposed as a simple and easy
496 measure of longitudinal function in healthy volunteers
497 and patients with hypertrophic cardiomyopathy (39).
498 Longitudinal contraction assessed in the form of mitral
499 annular plane systolic excursion (MAPSE) demonstrated
500 that despite similar LV ejection fraction of >55%, healthy
501 volunteers appeared to have a higher baseline MAPSE than
502 those patients with severe mitral regurgitation; reflecting a
503 better longitudinal contractility of the left ventricle. During
504 exercise at 55% HR_{max} , healthy volunteers had an improved
505 longitudinal contractility before the value plateaued at 75%
506 HR_{max} . This initial change of improved contractility was not
507 seen in patients with severe mitral regurgitation, potentially
508 reflecting a deconditioned myocardium. These results
509 should however be interpreted with caution in light of the
510 relatively small sample population.

511 Intra-observer reproducibility of LV parameters was
512 excellent at all three stages. Similarly, inter-observer
513 reproducibility of LV parameters was also excellent.
514 Although RVESV measurements were the least reproducible
515 during exercise, the RV ejection fraction and cardiac
516 index were however highly reproducible at all 3 stages.
517 The inter-scan reproducibility was less optimal for LV
518 parameters (CV 5–16%) and RVESV (CV 11–24%). The
519 wide interscan variability can possibly be explained by
520 the long 16 weeks scan interval between the 1st and 2nd
521 exCMR scans. Although healthy volunteers had no specific
522 exercise training during that period, other factors such as
523 different loading conditions, diet and temperatures could
524 influence cardiac physiology on a day-to-day basis.

525 This study has highlighted the potential of using
526 ‘navigated’ image acquisition techniques for the assessment
527 of cardiovascular response during continuous exercise.
528 ExCMR has the potential of providing quantitative cardiac

529 indices, whilst offering a direct link between physical
 530 activity, symptoms and stress imaging findings. Additionally,
 531 it can offer important information such as functional
 532 capacity and BP response. The use of exCMR can create
 533 new avenues for research and clinical practice, such as stress
 534 evaluation of ventricular dysfunction. This is particularly
 535 relevant to pathologies of the LV and RV, and pulmonary
 536 circulation that are challenging to assess by other imaging
 537 modalities. Further assessment of this ex-CMR protocol is
 538 now warranted for assessment of cardiac pathologies where
 539 current exercise imaging modalities have been shown to
 540 have limitations.

541

542 *Limitations*

543 As per all supine exCMR studies, there are general
 544 limitations with this approach. Cycling whilst lying in a
 545 flat, supine position is an unorthodox form of exercise,
 546 and skeletal muscle fatigue may lead to premature test
 547 termination (20). Knee-to-bore clearance whilst cycling is
 548 also limited by patient height and magnet bore diameter.
 549 This study had a maximum participant's height of 188 cm.
 550 Furthermore, vigorous respiratory movement can also
 551 result in blurring or ghosting of images collated across
 552 cardiac cycles. When respiration is performed in the
 553 anterior-posterior direction, thus not captured by the
 554 navigator in the head-feet direction, the navigator could
 555 potentially fail to work. The use of respiratory navigator
 556 also causes interruption to steady-state imaging and these
 557 signal variations can potentially lead to artifacts, particularly
 558 in the systole phase during exercise. Optimal patient
 559 preparation, as detailed in the methodology, is therefore
 560 vital. Other limitations of exCMR include its inability to be
 561 performed in patients with certain implanted devices. Since
 562 most CMR acquisitions are acquired over multiple cardiac
 563 cycles, arrhythmias such as atrial fibrillation or premature
 564 ventricular contractions may pose additional challenges
 565 for standard CMR sequences. The study population
 566 was small, and the reproducibility should therefore be
 567 interpreted with caution. Although highly reproducible,
 568 the findings of this study were also not validated against
 569 an invasive reference standard. Further work could look
 570 into assessing the accuracy of this imaging method against
 571 invasive exercise standards (direct Fick method) in deriving
 572 cardiac output. This technique was not intended to achieve
 573 the 85% of 'age-predicted maximal heart rate' required
 574 for myocardial ischaemia testing purposes, as in-scanner
 575

12 lead ECG monitoring is not feasible, and therefore
 accurate assessment of ST segment changes during exercise,
 which may prompt test termination, cannot be performed.
 The primary aim of this navigated exCMR technique was
 to assess the serial change in ventricular volumes with
 exercise as this can serve as an important tool in enabling
 understanding of physiology in patients with exertional
 symptoms and structural/congenital heart disease.

Conclusions

This exercise CMR protocol using a novel application of
 the free-breathing, multi-shot, navigated imaging method
 allows simultaneous assessment of the left and RV volumes
 during *continuous* exercise. This study demonstrates
 feasibility of exCMR in patients with mitral regurgitation
 for the first time. Intra and inter-observer readings were
 highly reproducible. Clinical feasibility of this protocol
 suggests a future role in the assessment of patients with
 exercise-related symptoms.

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 YH/0551) and complied with the Declaration of Helsinki
 (as revised in 2013). All participants provided written

624 informed consent.

625

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4 Chamber exCMR cine images

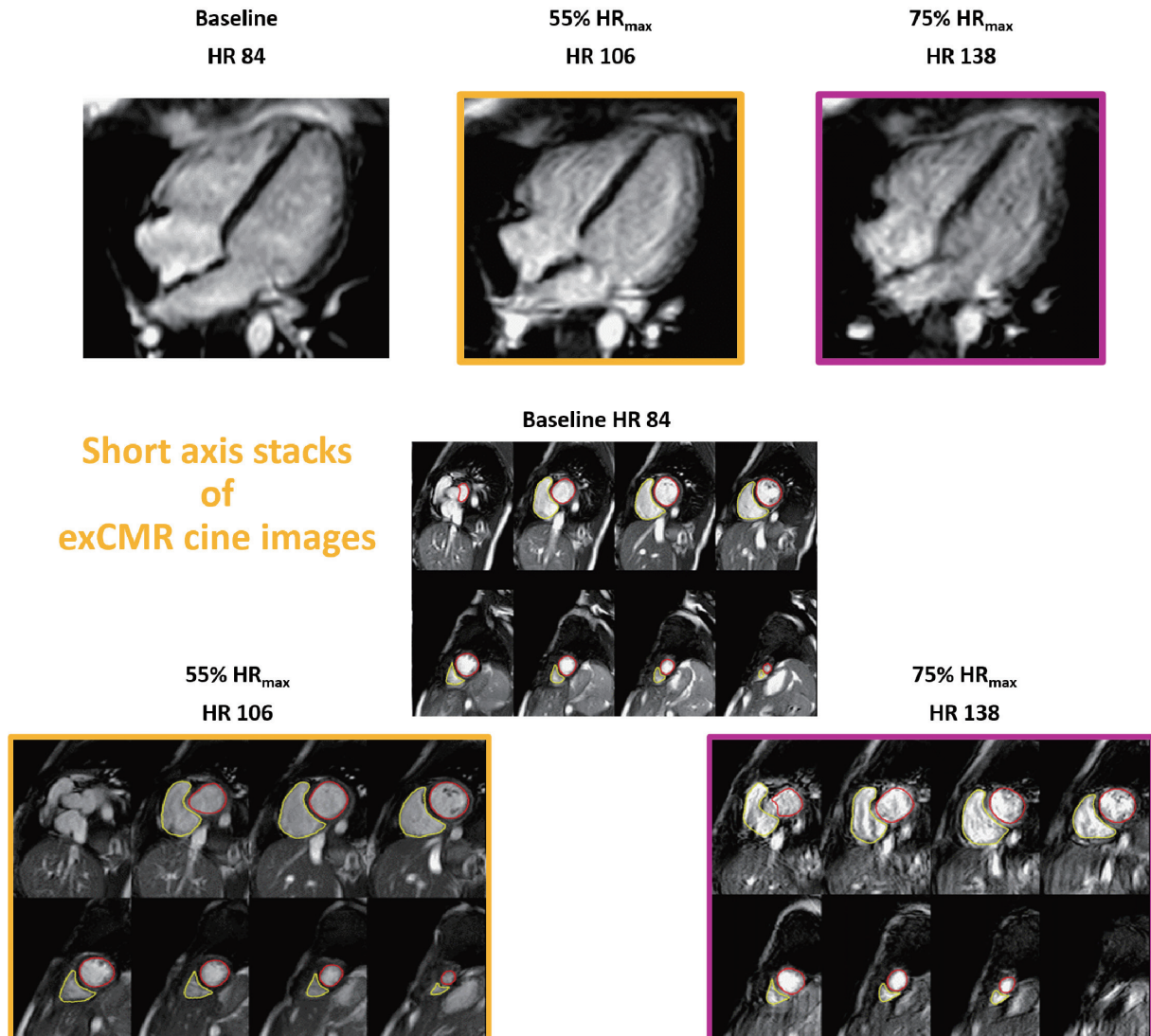


Figure S1 Illustration of navigated exCMR cine images from one volunteer. Cine images at baseline, 55% HR_{max} and 75% HR_{max}. Heart rate at time of imaging was 84, 106 and 138 bpm, respectively.