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ORIGINAL RESEARCH ARTICLE

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Is Cardiorespiratory Fitness Related to Cardiometabolic Health and All-Cause Mortality Risk in Patients with Coronary Heart Disease? A CARE CR Study

Simon Nichols^{1*} , Claire Taylor², Richard Page³, Anna Kallvikbacka-Bennett⁴, Fiona Nation³, Toni Goodman⁵, Andrew L. Clark⁴, Sean Carroll³ and Lee Ingle³

Abstract

Background: Higher cardiorespiratory fitness (CRF) is associated with lower morbidity and mortality in patients with coronary heart disease (CHD). The mechanisms for this are not fully understood. A more favourable cardiometabolic risk factor profile may be responsible; however, few studies have comprehensively evaluated cardiometabolic risk factors in relation to CRF amongst patients with CHD. We aimed to explore differences in cardiometabolic risk and 5-year all-cause mortality risk in patients with CHD who have low, moderate, and high levels of CRF.

Methods: Patients with CHD underwent maximal cardiopulmonary exercise testing, echocardiogram, carotid intima-media thickness measurement, spirometry, and dual X-ray absorptiometry assessment. Full blood count, biochemical lipid profiles, high-sensitivity (hs) C-reactive protein, and NT-proBNP were analysed. Patients were defined as having low, moderate, or high CRF based on established prognostic thresholds.

Results: Seventy patients with CHD (age 63.1 ± 10.0 years, 86% male) were recruited. Patients with low CRF had a lower ventilatory anaerobic threshold, peak oxygen pulse, post-exercise heart rate recovery, and poor ventilatory efficiency. The low CRF group also had higher NT pro-BNP, hs-CRP, non-fasting glucose concentrations, and lower haemoglobin and haematocrit. Five-year mortality risk (CALIBER risk score) was also greatest in the lowest CRF group (14.9%).

Conclusions: Practitioners should interpret low CRF as an important clinical risk factor associated with adverse cardiometabolic health and poor prognosis, study registry; www.researchregistry.com.

Keywords: Coronary Heart Disease, Cardiac Rehabilitation, Cardiometabolic Health, Exercise Training, Atherosclerosis, VO_{2peak} , Maximal Cardiopulmonary Exercise Testing, CALIBER 5-year risk

Key Points

1. Low cardiorespiratory fitness is associated with the poorest cardiometabolic health in patients with coronary heart disease.
2. Five-year risk of all-cause death and NT-proBNP are highest amongst coronary heart disease patients

with the lowest cardiorespiratory fitness, even when left ventricular ejection fraction is preserved. Longer-term or higher intensity exercise-based cardiac rehabilitation programmes that closely monitor cardiovascular risk factors may be warranted for coronary heart disease patients who have low cardiorespiratory fitness.

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Background

Cardiorespiratory fitness (CRF) or VO_{2peak} predicts all-cause and cardiovascular (CV) mortality in patients

with coronary heart disease (CHD) [1–3]. We have shown that in patients with CHD, low CRF confers the highest mortality risk (54%) over a 14-year period when compared to those categorised as having moderate (31%) or high CRF [17%] [4]. Improvements in CRF resulting from exercise-based cardiac rehabilitation (CR) may reduce mortality in patients who have low CRF [4–6].

Whilst higher CRF is a strong predictor of better survival outcomes, the mechanisms responsible for this association are not fully understood. Few studies have comprehensively profiled the cardiometabolic health status of patients with CHD in relation to established CRF categories. Higher all-cause mortality in patients with CHD may not be entirely attributable to older age, cardiovascular disease (CVD) severity, and more comorbidities [7]. Unidentified and potentially treatable cardiometabolic risk factors may cause the divergence in mortality rates observed between patients with high and low CRF.

Biomarkers of cardiac dysfunction, inflammation, cardiac autonomic function, arterial plaque status, renal function, blood oxygen carrying capacity, and metabolic control have received little attention in relation to CRF in patients with CHD. Where limited data does exist, CRF is often reported as estimated metabolic equivalents (METs) from treadmill or cycle ergometer workloads [2, 4, 6]. Fitness estimation may be inaccurate in patients with CHD, could result in individual patients being inappropriately assigned to a specific CRF category and, attenuate any prognostic signal [8]. The association between cardiometabolic health and CRF should be investigated using ‘gold-standard’ maximal cardiopulmonary exercise testing (CPET).

This cross-sectional investigation aimed to assess differences in CV and cardiometabolic health amongst patients with CHD when characterised as have low, moderate, or high levels of CRF. We also investigated differences in estimated 5-year risk of death using the validated CALIBER composite scoring system [9].

Methods

Study Design

Data for this study were taken from the Cardiovascular and cardiorespiratory Adaptations to Routine Exercise-based Cardiac Rehabilitation study (CARE CR). Ethical approval for the study was given by the Humber Bridge NHS Research Ethics Committee-Yorkshire and the Humber (12/YH/0278). The study is registered with www.researchregistry.com (researchregistry3548). All procedures were conducted in accordance with the ethical standards outlined in the 1964 Helsinki declaration and its later amendments.

The methodology for this study has been reported elsewhere [10]. In brief, patients were recruited following a recent hospital admission and referral to CR for stable

angina, myocardial infarction (MI), coronary artery bypass graft (CABG) surgery, or elective percutaneous coronary intervention (PCI). After giving verbal consent, patients were invited for assessment at Academic Cardiology, Castle Hill Hospital, Kingston-Upon-Hull, United Kingdom, where written informed consent was obtained.

Resting Haemodynamics, Anthropometry, and Body Composition

Resting heart rate (HR) was determined at the end of 15 min, semi-supine rest using a 12-lead ECG (GE Healthcare, Buckinghamshire, UK). Left arm brachial blood pressure was recorded using an ECG-gated automated blood pressure (BP) cuff (Tango, SunTech Medical, Eynsham, UK).

Stature (cm) was measured using a Leicester Height Measure (SECA, Birmingham, UK) with patients standing, without footwear, in the Frankfort plane with their heels and head positioned to the back of the stadiometer. Waist circumference measurements were taken 1 cm above the iliac crest [11], and hip measurements were taken from the widest aspect of the buttocks using an inflexible tape measure. Both measurements were recorded in centimetre, and a ratio of the two was calculated to determine waist to hip circumference ratio [11].

Body composition was analysed using dual X-ray absorptometry [DXA] (Lunar iDXA, GE Healthcare, Buckinghamshire, UK). Total body mass (kg), lean body mass (kg), total fat (%), android fat (%), and android gynoid ratio were determined using the Lunar iDXA's integrated software. DXA-derived total body mass was used to determine body mass index (BMI; $\text{kg}\cdot\text{m}^{-2}$).

Echocardiogram

Standard 2D, M-mode echocardiogram techniques were used to determine left ventricular (LV) function. LV ejection fraction (LVEF) was calculated using the Simpson's method from measurements of end-diastolic and end-systolic volumes on apical 4-chamber and 2-chamber 2D views following the guidelines of Schiller and colleagues [12]. LV systolic dysfunction was diagnosed if LVEF was $\leq 45\%$.

Carotid Intima-Media Thickness and Carotid Plaque Measurement

Carotid intima-media thickness (C-IMT) and carotid plaque measurements were obtained using the Panasonic CardioHealth Station (Panasonic Biomedical Sales Europe BV, Leicestershire, UK) which has low measurement variability [13, 14]. Measurements were taken using previously outlined methods [10, 13] from a 1 cm segment of the common carotid artery (CCA) located 1 cm proximally from the carotid bifurcation. Measurements from the right and left CCA were taken in the longitudinal plane at anterior (right CCA 150°; left CCA

210°), lateral (right CCA 120°; 230°), and posterior (right CCA 90°; left CCA 270°) angles, relative to ground. Each C-IMT measurement was determined from the average of 24 digital distance markers automatically placed between the intimal and medial boundaries and reported to three decimal places. Carotid artery plaques were measured in the longitudinal plane. The largest plaque was taken as the representative measure of the patient's carotid plaque status and reported as being either < 1.0 mm, ≥ 1.0 to < 3.0 mm or ≥ 3.00 mm.

Blood Samples

Blood samples were taken from a vein at the antecubital fossa and collected in ethylenediaminetetraacetic acid (EDTA), potassium oxalate, and serum separating tubes (SST). Full blood count, (haematocrit, haemoglobin, and neutrophil and lymphocyte count) and estimated glomerular filtration rate (eGFR) were analysed on the day of collection using a registered National Health Service (NHS) pathology lab (Castle Hill Hospital, Hull, United Kingdom). A further EDTA and potassium oxalate tube were placed in a refrigerated (4 °C) centrifuge at 3000 rpm, for 15 min immediately after the blood draw. Samples collected in SST tubes were allowed to clot for 30 min prior to being centrifuged under the same conditions.

The ABX Pentra 400 biochemistry auto analyser (Horiba, Montpellier, France) was used to analyse serum triglycerides, total cholesterol, high-density lipoprotein cholesterol (HDL), plasma glucose, and high-sensitivity C-reactive protein (hs-CRP) in duplicate. Calibration and quality controls were conducted in accordance with manufacturer's guidelines. Low-density lipoprotein (LDL) was estimated using the Friedewald equation [15].

Cardiopulmonary Exercise Testing and Assessment of Physical Activity

After 3 min of seated rest, CPET was conducted following the modified Bruce treadmill protocol (GE Healthcare, Buckinghamshire, UK) [16] in accordance with established guidelines [17–20]. A 12-lead ECG was monitored continuously throughout the test. An ECG-gated automated BP measurement was recorded at the start of CPET and at the second minute of a test stage until the end of the test. RPE scores were recorded at peak exercise. HR was recorded at 1, 2, 3, and 6 min during the passive seated recovery period.

Breath-by-breath gas exchange data were collected using an Oxycon Pro metabolic cart (Jaeger, Hoechburg, Germany). VO_{2peak} was defined as the mean VO_2 (ml) over the last 30 s of the CPET. VO_{2peak} was adjusted for both body mass and lean (DXA-derived) body mass (ml/kg/min). The ventilatory anaerobic threshold (VAT) was determined using

the V-slope method [21] by two independent investigators. Where agreement on VAT scores was not met, the results were discussed with a third independent investigator and a consensus was reached. The VAT was determined using the average of the middle 5 of 7 consecutive breaths (excluding the highest and lowest measures) and reported standardised to patient body mass (ml/kg/min). The oxygen uptake efficiency slope (OUES), VE/VCO_2 slope, and oxygen pulse (VO_2/HR) were calculated as previously described [19]. Self-reported weekly physical activity levels were obtained by asking patients if they participated in either 150 min of moderate physical activity, 75 min of vigorous physical activity, or both.

Determination of Cardiorespiratory Fitness Categories

Patients were categorised into three groups based on their VO_{2peak} . We converted VO_{2peak} into metabolic equivalents (METs) by dividing by 1 MET (estimated to be equivalent to a VO_2 of 3.5 ml/kg/min) to allow for easier comparisons with other studies. Low CRF was defined as a peak MET value of < 5 for men and < 4 for women; high CRF was defined as peak MET > 7 for men and > 6 for women. These thresholds were based on previously published prognostic thresholds [4–6, 22, 23].

Prognosis—CALIBER 5-year Risk Score

A 5-year risk of all-cause mortality was calculated for each patient using the comprehensive online (www.caliberresearch.org/model) CALIBER 5-year risk score [9]. This model does not include any CRF measurements in its calculation. A full list of included variables can be found in Table 1. Five-year risk of all-cause mortality was reported as a percentage to one decimal place.

Statistical Analysis

Statistical analysis was performed using SPSS version 22 (IBM, New York, USA). The Shapiro-Wilk test and histograms were used to assess normality. Categorical data are reported as percentages. Continuous normally distributed variables are displayed as mean with 95% confidence intervals (95% CI) or standard deviation (\pm) where specified. One-way analysis of variance (ANOVA), one-way analysis of covariance (ANCOVA), and chi-squared analysis were used to assess differences between CRF groups. The only covariate considered in analysis was age as sex was used in assigning patients to their CRF category.

Statistical significance was set at $p < 0.05$. Partial eta squared (η_p^2) effect sizes were used to report the magnitude of group differences. Effect sizes of 0.01, 0.06, and 0.14 denoted small, moderate, and large effect sizes, respectively [24]. Pearson correlations were used to assess the strength of the relationship between CRF category

Table 1 Variables Included in the CALIBER 5-Year Risk Score

| Categorical Variables | Continuous Variables |
|-------------------------------------|-------------------------------|
| Sex | Age (years) |
| Belongs to most deprived quintile | Total cholesterol (mmol/L) |
| CAD diagnosis and severity | HDL (mmol/L) |
| Interventions (last 6 months) | Heart rate (beats per minute) |
| Smoking status | Creatinine (micromol/L) |
| Hypertension/BP lowering medication | White cell count ($10^9/L$) |
| Diabetes | Haemoglobin (g/dl) |
| Heart failure | |
| Peripheral arterial disease | |
| Atrial fibrillation | |
| Stroke | |
| Chronic renal disease | |
| COPD | |
| Cancer | |
| Chronic liver disease | |
| Depression | |
| Anxiety | |

CAD coronary artery disease, BP blood pressure, COPD chronic obstructive pulmonary disease, HDL high-density lipoprotein

and CALIBER 5-year risk of death. An r value of < 0.25 , 0.26 to 0.50, 0.51 to 0.75, and, > 0.75 were considered weak, moderate, fair, and strong associations, respectively [25].

Results

Cohort Characteristics

Seventy patients were recruited. Sixteen (23%) patients had sustained an ST-elevation myocardial infarction (STEMI), 22 (31%) had sustained a non-STEMI, and 19 (27%) had undergone elective PCI. Seven patients (10%) were medically managed for stable angina, and six (9%) had undergone CABG. The mean age of the cohort was 63.1 ± 10.0 years (BMI $29.2 \text{ kg m}^{-2} \pm 4.0$; 86% male). The mean LVEF was $55.0 \pm 6.9\%$. The median time from cardiac event to consent was 54 days (range 22 to 220). 94% of patients were seen within 90 days.

Twenty-eight (40%), 32 (46%), and 10 (14%) of patients were defined as having high, moderate, and low levels of CRF, respectively. CRF group characteristics are shown in Table 2. Patients with low CRF were older ($p < 0.001$) and had a lower lean body mass than patients with moderate ($p = 0.029$) and high CRF ($p = 0.002$). Compared to patients with high ($p < 0.001$) and moderate CRF ($p = 0.002$), patients with low CRF also had a lower android gynoid ratio, higher resting HR (high CRF, $p = 0.004$; moderate CRF, $p = 0.042$), and a larger proportion of patients suffering from type II diabetes ($p = 0.028$). Medications (Table 3) were comparable across all

three groups with the exception of diuretics which were more commonly prescribed to patients in the low CRF group ($p = 0.001$).

Cardiorespiratory Fitness and Physical Activity

Cardiorespiratory fitness and physical activity-related variables are shown in Table 4. Differences in $VO_{2\text{peak}}$ standardised to lean body mass, VAT, VO_2/HR , and exercise test duration were observed across all groups. Compared to patients with high CRF, VE/VCO_2 slope and OUES were both higher amongst patients in the moderate and low CRF groups. Mean LVEF was not different across the groups and was not significantly correlated with peak METs ($r = 0.147$; $p = 0.224$). One patient in both the low and moderate CRF groups had a LVEF $< 40\%$. Compared to patients with high and moderate CRF, patients with low CRF had the most impaired 1-min HR recovery, an indicator of cardiac autonomic function. The proportion of patients who reported participating in either 150 min of moderate ($p = 0.011$) or 75 min of vigorous physical activity per week was higher in the high CRF group.

Blood Biomarkers and Cardiovascular Risk

Results of blood biochemical analyses are displayed in Table 5. NT-proBNP was inversely associated with CRF ($r = -0.414$) and was highest amongst patients with low CRF. Four (40%) patients with low CRF had an NT-proBNP > 400 pg/ml compared to eight (25%) and one (4%) in the moderate and high CRF groups. One patient in the low CRF group without previously diagnosed chronic heart failure (CHF) had an NT-proBNP result > 2000 pg/ml with a mildly reduced LVEF (45%), sinus rhythm, and with an eGFR of $73 \text{ mL/min/1.73 m}^2$. Lipid profiles did not differ between groups. Non-fasting plasma glucose concentrations in the low CRF group were higher than the moderate CRF group ($p = 0.008$). hs-CRP was also highest in the low CRF group. Haematocrit and haemoglobin differed between all groups.

Carotid Intima-Media Thickness

A moderate inverse correlation between left-sided C-IMT measurements and CRF was observed ($r = -0.382$; $p = 0.001$) with larger measurements seen in the low (0.926; 95% CI 0.797, 1.054; $p = 0.002$) and moderate CRF groups (0.827; 95% CI 0.755, 0.898; $p = 0.011$) compared to the high CRF group [0.689; 95% CI 0.612, 0.765] (Fig. 1). There was no significant correlation between right sided C-IMT and CRF ($r = -0.210$; $p = 0.080$). The proportion of patients with left-sided plaque score < 1 mm (Fig. 2) increased with each CRF group (low 10%; moderate 25%; high 39%). The proportion of patients exhibiting large left-sided plaque scores (> 3 mm) decreased across each group (low 40%; moderate 16%; high 4%; $p =$

Table 2 Patient Characteristics Expressed as Mean (95% Confidence Intervals)

| Variable | High CRF, n = 28 | Mod CRF, n = 32 | Low CRF, n = 10 | Partial eta squared | p value |
|-----------------------------|---------------------------------|--------------------------------|--------------------------------|---------------------|-----------------------|
| Sex (% male) | 26 (93) | 27 (84) | 7 (70) | | 0.199 |
| Age (years) | 56.3 (53.1, 59.4) ^{†*} | 67.2 (64.3, 70.2) [†] | 69.3 (64.1, 74.5) [*] | 0.326 | < 0.001 ^{**} |
| BMI (kg.m ⁻²) | 29.0 (27.5, 30.5) | 28.9 (27.5, 30.3) | 30.9 (28.3, 33.4) | 0.030 | 0.363 |
| Waist circumference (cm) | 101.3 (97.3, 105.4) | 102.6 (98.7, 106.4) | 108.1 (101.4, 114.9) | 0.043 | 0.235 |
| Waist to hip ratio (cm) | 0.96 (0.93, 0.98) | 0.98 (0.95, 1.00) | 0.96 (0.92, 1.01) | 0.018 | 0.553 |
| Android fat % | 45.0 (42.0, 48.0) | 48.0 (45.2, 50.8) | 47.9 (42.9, 52.9) | 0.034 | 0.316 |
| Total fat % | 35.9 (32.2, 39.6) | 37.3 (33.8, 40.7) | 41.1 (34.8, 47.3) | 0.029 | 0.371 |
| Lean body mass (Kg) | 55.0 (51.9, 58.1) ^{†*} | 50.2 (47.3, 53.1) [†] | 45.3 (40.1, 50.5) [*] | 0.145 | 0.005 ^{**} |
| Android/gynoid ratio | 1.31 (1.24, 1.38) [*] | 1.26 (1.20, 1.32) ^x | 1.06 (0.95, 1.17) ^x | 0.182 | 0.001 ^{**} |
| LVEF (%) | 56.6 (54.0, 59.2) | 54.3 (51.8, 56.6) | 52.8 (48.5, 57.1) | 0.043 | 0.232 |
| Resting HR (bpm) | 56 (52, 69) [*] | 59 (56, 63) ^x | 67 (61, 74) ^x | 0.118 | 0.015 ^{**} |
| Resting SBP (mmHg) | 130 (122, 137) | 124 (117, 131) | 138 (125, 150) | 0.056 | 0.147 |
| Resting DBP (mmHg) | 85 (80, 89) | 82 (77, 86) | 73 (65, 81) | 0.083 | 0.055 |
| FEV ₁ /FVC ratio | 0.78 (0.75, 0.81) | 0.75 (0.72, 0.78) | 0.75 (0.70, 0.80) | 0.042 | 0.237 |
| Presenting diagnosis | | | | | |
| MI (%) | 17 (61) | 15 (47) | 6 (60) | | 0.642 |
| PCI (%) | 8 (29) | 11 (34) | 1 (10) | | |
| Angina (%) | 2 (7) | 3 (9) | 2 (20) | | |
| CABG (%) | 1 (4) | 3 (9) | 1 (10) | | |
| Past medical history | | | | | |
| Previous MI (%) | 6 (21) | 6 (19) | 2 (20) | | 0.186 |
| Type II diabetes (%) | 3 (12) | 6 (19) | 5 (50) | | 0.028 ^{**} |
| Atrial fibrillation (%) | 0 (0) | 2 (6) | 1 (10) | | 0.269 |
| Smoker (%) | 1 (4) | 1 (3) | 2 (20) | | 0.238 |
| Ex-smoker (%) | 16 (57) | 16 (50) | 7 (70) | | |

Mod moderate, *BMI* body mass index, *LVEF* left ventricular ejection fraction, *HR* heart rate, *SBP* systolic blood pressure, *DBP* diastolic blood pressure, *FEV1/FVC* ratio of forced expiratory volume in 1 s to forced vital capacity, *MI* myocardial infarction, *PCI* percutaneous coronary intervention, *CABG* coronary artery bypass graft

^{**}Significant group effect

^{*}Significant difference between high CRF and low CRF

[†]Significant difference between high CRF and moderate CRF

^xSignificant difference between mod CRF and low CRF

0.047). Patients with moderate and low CRF had incrementally higher proportions of patients with right-sided plaque scores > 3 mm (moderate CRF 9.4%; low CRF 30%) compared to patients in the high CRF who had no plaques > 3 mm (Fig. 2; $p = 0.032$).

All-Cause Mortality Estimation

Differences in CALIBER 5-year risk (Fig. 3) were observed between all groups. Estimated risk was highest in the low CRF group (14.9; 95% CI 11.4, 18.5%). The moderate group had a 9.7% 5-year risk of death (95% CI 7.7, 11.7%) whilst high CRF was associated with lowest 5-year risk of death (3.7%; 95% CI 1.6, 5.8%). There was a negative association between peak METS and CALIBER 5-year risk score ($r = -0.538$; $r^2 = 0.289$; $p < 0.001$).

Statistical Adjustment for Age

Differences in left-sided mean C-IMT ($p = 0.274$; $\eta_p^2 = 0.039$) became non-significant following age adjustment. Statistically significant differences for all other variables remained unaltered after statistical adjustment for age.

Discussion

To our knowledge, this is the first study to comprehensively profile the cardiometabolic and CV health status of patients with CHD according to prognostically verified CRF categories [4–6, 22, 23]. Patients with the lowest CRF have the poorest integrated cardiorespiratory function and cardiometabolic health. Patients with low CRF may also have impaired autonomic function and worse CV disease severity. Low CRF was associated with the poorest CALIBER 5-year all-cause mortality risk and 'high risk' status (defined as 3%

Table 3 Number (%) of Medications Taken by Patients

| Medication | High CRF, <i>n</i> = 28 | Mod CRF, <i>n</i> = 32 | Low CRF, <i>n</i> = 10 | <i>p</i> value |
|-----------------------------------|-------------------------|------------------------|------------------------|----------------|
| Aspirin (%) | 27 (96) | 31 (97) | 10 (100) | 0.838 |
| Ticagrelor (%) | 15 (54) | 17 (53) | 3 (30) | 0.393 |
| Clopidogrel (%) | 10 (36) | 8 (25) | 3 (30) | 0.665 |
| Anti-coagulants (%) | 0 (0) | 1 (3) | 1 (10) | 0.263 |
| Beta-blockers (%) | 24 (86) | 29 (91) | 7 (70) | 0.266 |
| ACE-inhibitors (%) | 19 (68) | 16 (50) | 7 (70) | 0.291 |
| Angiotensin receptor blockers (%) | 0 (0) | 5 (16) | 1 (10) | 0.096 |
| Statins (%) | 25 (89) | 32 (100) | 10 (100) | 0.095 |
| Diuretics (%) | 1 (4) | 1 (3) | 5 (50) | 0.001** |
| Calcium channel blockers (%) | 4 (14) | 8 (25) | 4 (40) | 0.502 |
| Nitrates (%) | 2 (7) | 3 (9) | 3 (30) | 0.233 |

ACE angiotensin converting enzyme

**Significant group effect

Table 4 Cardiorespiratory Fitness and Physical Activity Characteristics Expressed as Mean (95% Confidence Intervals)

| Variable | High CRF, <i>n</i> = 28 | Mod CRF, <i>n</i> = 32 | Low CRF, <i>n</i> = 10 | Partial eta squared |
|--|---------------------------------------|--------------------------------------|--------------------------------------|---------------------|
| VO _{2peak} (ml/kg/min) | 28.5 (27.3, 29.7) ⁺ | 20.7 (19.5, 21.8) ⁺ | 14.9 (12.8, 16.9) ⁺ | – |
| VO _{2peak} (L/min) | 2478.2 (2333.0, 2623.5) ⁺ | 1749.0 (1613.1, 1884.9) ⁺ | 1273.8 (1030.7, 1516.8) ⁺ | – |
| VO _{2peak} -lean (ml/kg/min) | 45.2 (43.4, 47.0) ⁺ | 34.8 (33.2, 36.5) ⁺ | 26.8 (23.8, 29.8) ⁺ | 0.670 |
| VAT (ml/kg/min) | 20.7 (19.3, 22.1) ⁺ | 14.6 (13.3, 15.9) ⁺ | 11.2 (8.9, 13.6) ⁺ | 0.494 |
| VE/VCO ₂ slope | 30.1 (28.2, 32.1) ^{*†} | 37.4 (35.6, 39.2) [†] | 38.5 (35.2, 41.7) [*] | 0.354 |
| VO ₂ /HR (ml/beat) | 17.0 (15.8, 18.2) ⁺ | 13.8 (12.7, 14.9) ⁺ | 11.3 (9.4, 13.3) ⁺ | 0.311 |
| OUES | 2718.3 (2555.3, 2881.3) ^{*†} | 1963.5 (1811.1, 2116.0) [†] | 1699.0 (1426.2, 1971.7) [†] | 0.485 |
| eBR (%) | 30.3 (23.6, 36.9) | 28.1 (22.0, 34.3) | 37.0 (26.0, 48.1) | 0.028 |
| Peak HR (bpm) | 147 (141, 153) ^{*†} | 128 (122, 134) [†] | 119 (108, 129) [*] | 0.308 |
| Peak RER | 1.13 (1.09, 1.12) [*] | 1.09 (1.05, 1.12) ^x | 0.97 (0.91, 1.04) ^{**x} | 0.181 |
| Peak RPE | 18 (17, 19) | 18 (17, 19) | 17 (15, 18) | 0.072 |
| 1 min HR recovery (bpm) | –36 (–32, –40) [†] | –30 (–26, –34) ^x | –18 (–11, –25) ^{**x} | 0.209 |
| 2 min HR recovery (bpm) | –54 (–50, –59) ⁺ | –45 (–40, –49) ⁺ | –32 (–25, –38) ⁺ | 0.312 |
| 3 min HR recovery (bpm) | –60 (–56, –65) ⁺ | –49 (–45, –53) ⁺ | –37 (–30, –44) ⁺ | 0.359 |
| 6 min HR recovery (bpm) | –67 (–62, –71) ⁺ | –54 (–50, –58) ⁺ | –41 (–33, –48) ⁺ | 0.377 |
| Exercise test duration (s) | 963.2 (916.3, 1010.1) ⁺ | 747.8 (703.9, 791.6) ⁺ | 488.3 (409.8, 566.8) ⁺ | 0.635 |
| METs | 8.1 (7.8, 8.5) ⁺ | 5.9 (5.6, 6.2) ⁺ | 4.3 (3.7, 4.8) ⁺ | – |
| Maximal CPET (%) | 26 (93) | 26 (81) | 6 (60) | 0.058 |
| Achieves 150 min of moderate activity per week (%) | 18 (64) ⁺ | 9 (28) | 5 (50) | 0.011 ^{**} |
| Achieves 75 min of vigorous activity per week (%) | 7 (25) ⁺ | 1 (3) | 0 (0) | 0.013 ^{**} |

VO_{2peak} peak oxygen uptake, VAT ventilatory anaerobic threshold, VE/VCO₂ ventilatory efficiency with respect to CO₂ elimination, VO₂/HR oxygen pulse, OUES oxygen uptake efficiency slope, eBR estimated breathing reserve, HR heart rate, bpm beats per minute, RER respiratory exchange ratio, RPE rating of perceived exertion; S seconds, METs metabolic equivalents

**Significant group effect

*Significant difference between high CRF and low CRF

†Significant difference between high CRF and moderate CRF

xSignificant difference between mod CRF and low CRF

+Significantly different from all other groups

Table 5 Blood Biomarkers Expressed as Mean (95% Confidence Intervals)

| Variable | High CRF, n = 28 | Mod CRF, n = 32 | Low CRF, n = 10 | Partial eta squared | p value |
|------------------------------------|-----------------------------------|-----------------------------------|-----------------------------------|---------------------|-----------|
| NT-proBNP (ng/L) | 86.1 (11.4, 1428.0) *† | 217.0 (33.9, 1916.0)† | 464.0 (26.9, 2735.0)* | - | < 0.001** |
| Total Chol (mmol/L) | 3.6 (3.2, 3.9) | 3.7 (3.4, 4.0) | 3.9 (3.3, 4.5) | 0.017 | 0.579 |
| LDL Chol (mmol/L) | 1.7 (1.4, 1.9) | 1.8 (1.6, 2.1) | 1.9 (1.4, 2.3) | 0.019 | 0.529 |
| HDL Chol (mmol/L) | 1.2 (1.0, 1.3) | 1.2 (1.0, 1.3) | 1.4 (1.2, 1.6) | 0.052 | 0.176 |
| TC/HDL ratio | 3.2 (2.9, 3.5) | 3.3 (3.0, 3.6) | 3.0 (2.5, 3.6) | 0.01 | 0.713 |
| Triglycerides (mmol/L) | 1.6 (1.3, 1.9) | 1.5 (1.2, 1.8) | 1.4 (0.9, 2.0) | 0.011 | 0.691 |
| Glucose (mmol/L) | 5.7 (3.3, 13.8) | 5.2 (4.1, 22.2) ^x | 6.4 (5.5, 16.9) ^x | - | 0.012** |
| hs-CRP (mg/L) | 1.4 (0.3, 2.5)* | 2.8 (1.7, 3.8) | 4.2 (2.3, 6.2)* | 0.099 | 0.033** |
| Neutrophil/lymphocyte ratio | 2.4 (2.1, 2.8) | 2.8 (2.4, 3.1) | 3.3 (2.7, 3.8) | 0.084 | 0.052 |
| Haematocrit (%) | 0.423 (0.410, 0.435) ⁺ | 0.405 (0.393, 0.417) ⁺ | 0.376 (0.355, 0.397) ⁺ | 0.180 | 0.001** |
| Haemoglobin (g/L) | 145.5 (140.7, 150.2) ⁺ | 138.2 (133.7, 142.6) ⁺ | 125.6 (117.6, 133.6) ⁺ | 0.219 | < 0.001** |
| eGFR (mL/min/1.73 m ²) | 80.2 (75.2, 85.3) | 76.2 (71.4, 80.9) | 71.0 (62.6, 79.4) | 0.053 | 0.160 |

NT-proBNP N-terminal prohormone brain natriuretic peptides, Chol cholesterol, LDL low-density lipoproteins, HDL high-density lipoproteins, hs-CRP high-sensitivity C-reactive protein, eGF estimated glomerular filtration rate

**Significant group effect

*Significant difference between high CRF and low CRF

†Significant difference between high CRF and moderate CRF

^xSignificant difference between moderate CRF and low CRF

⁺Significantly different from all other groups

annual mortality risk [26]) despite the risk model not including CRF-based measurements. The CALIBER model includes an extensive set of all-cause mortality predictor variables, including sociodemographics, CHD severity and phenotype, primary CVD risk factors, CVD and non-CVD comorbidities, psychosocial factors, and biomarkers. This provides further evidence that CRF and other CPET-derived variables should be treated as ‘clinical vital sign’ [27].

An important finding of this study was that NT-proBNP levels and VE/VCO₂ slopes were highest in

the moderate and low CRF groups. This suggests a greater degree of cardiac dysfunction, ventilatory perfusion mismatch, or heightened peripheral chemoreceptor sensitivity as seen in patients with CHF [28, 29]. NT-proBNP levels and VE/VCO₂ slopes were higher even in the absence of significant resting LV impairment. It is possible that patients with CHD and low CRF have undiagnosed CHF or are in the early stages of its development. Future research is required to determine whether lower CRF in patients with CHD confers a

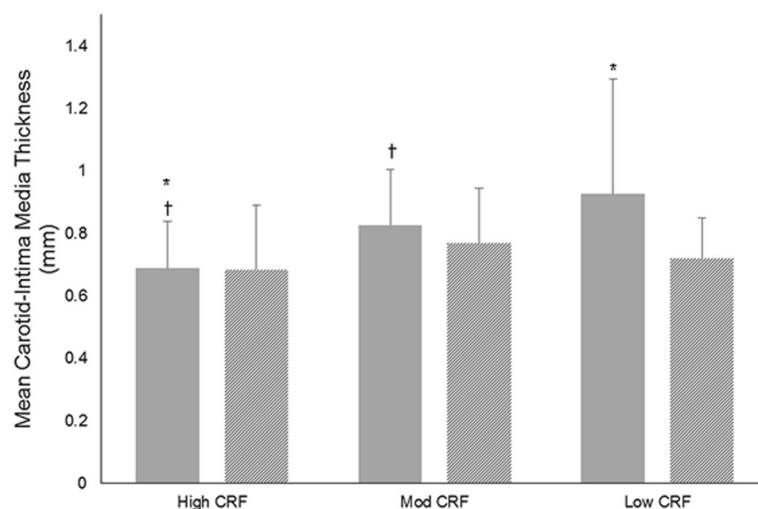


Fig. 1 Carotid intima-media thickness measurements in low, moderate, and high cardiorespiratory fitness groups. Mean left-sided carotid intima-media thickness measurements (solid grey bars) were higher in the low and moderate cardiorespiratory fitness categories. Mean right-sided carotid intima-media thickness (lines) did not differ between groups. Mod = moderate; CRF = cardiorespiratory fitness *Significant difference between high CRF and low CRF; †Significant difference between high CRF and moderate CRF

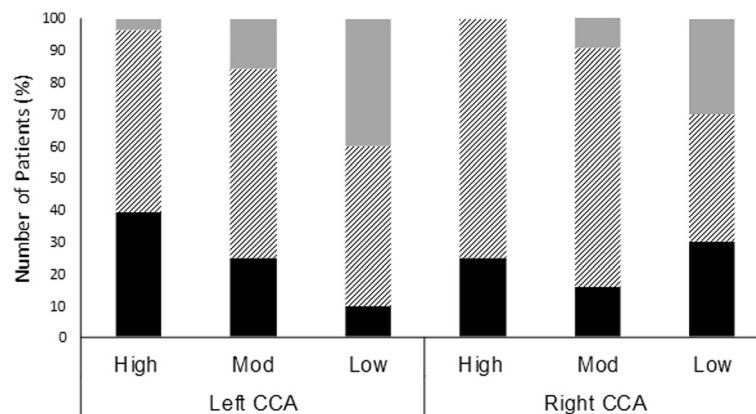


Fig. 2 Left- and right-side common carotid artery plaque severity in patients with high, moderate, and low cardiorespiratory fitness. Black bars indicate the proportion of patients with plaques < 1 mm and diagonal lines indicate the proportion of patients with plaques between 1 and 3 mm. Grey bars indicate the proportion of patients with plaques > 3 mm. Mod = moderate; CCA = common carotid artery

greater risk of developing CHF. Closer monitoring for signs and symptoms of CHF may be needed for patients who enter CR and have low CRF.

The prognostic value of a high VE/VCO₂ slope and/or NT-proBNP has been well-described in CHF [30–32]. However, the value of VE/VCO₂ slope and NT-proBNP levels has also been shown in CHD. Elevated values predict all-cause mortality [33], cardiovascular mortality (~5 year follow-up), and the development of CHF and stroke [34]. A high VE/VCO₂ is independently associated with high NT-proBNP levels, as well as ventricular remodelling [35]. Our findings suggest that patients with both low and moderate CRF levels are at greatest risk of having such adverse health outcomes, corroborated by their higher CALIBER 5-year risk.

Traditional risk factors such as angina symptoms, prior MI, atrial fibrillation, claudication, type II diabetes,

CHF, and smoking history do not entirely explain the increased mortality risk associated with having low CRF. Physical activity participation and higher CRF have been established as important independent causative factors for recurrent events [7]. In our study, patients defined as having high CRF reported participating in regular physical activity and had the lowest 5-year risk of death. This was accompanied by a lower resting HR and a quicker post-exercise HR recovery, indicators of autonomic function which also carry prognostic value. Interestingly however, no differences were evident in standard anthropometric indices (BMI or waist circumference) or in DXA-derived body fat content across the CRF groups.

The link between cardiometabolic profile, body composition, and cardiovascular risk has been widely reported and there is often an emphasis on reducing overweight and obesity states in CR programmes. In

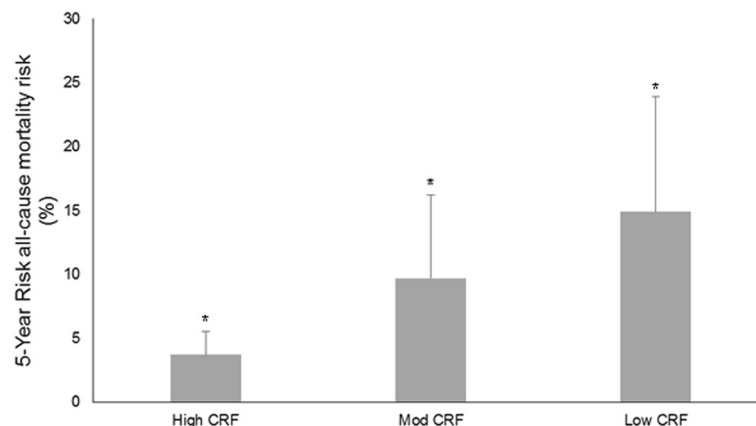


Fig. 3 CALIBER 5-year all-cause mortality risk scores were incrementally higher across the three cardiorespiratory fitness groups. Mod = moderate; CRF = cardiorespiratory fitness+Significantly different from all other groups

our study, patients with higher CRF had a larger proportion of lean tissue and more favourable abdominal fat distribution. Low CRF was only associated with latent changes in body composition. Changes were not evident when assessing patients with routinely applied anthropometric techniques, such as BMI and waist circumference. Whilst overweight patients with CHD have been reported to benefit from a better prognosis than their leaner counterparts, [36], the association between higher BMI and reduced all-cause mortality may depend on a patients' CRF. Those characterised as having low CRF appear to benefit from improved survival whilst fitter individuals do not [37, 38]. Our findings (incorporating criterion fitness and 'fatness' measures) suggest that maintaining or improving CRF may help lower recurrent cardiovascular risk in patients with CHD. However, our sample is relatively small and only 10 patients were identified as having a low CRF ($n = 10$). This may increase the likelihood of statistical error (type I/type II error) when comparing CRF groups. However, many of the effect sizes (η^2) associated with differences in cardiometabolic risk were large. Furthermore, the 95% CI indicate distinct values for many cardiometabolic variables across the CRF groups. However, further research using a larger patient cohort is required to confirm if our findings are representative of the wider CHD population.

As reported by others [4, 6], our data shows that patients with high CRF have lower CALIBER 5-year risk scores compared to patients with low and moderate CRF. In previous studies [4, 6], patients characterised as having high CRF were shown to have superior survival over a 15-year period. Barons and colleagues [6] reported that within their low CRF group, a history of MI, PCI, and angina were associated with the highest risk of death. Our data do not confirm this. However, similar to our findings, Taylor et al. [4] reported that prior MI is not more prevalent amongst patients with low CRF. It is possible that size and location of MI are responsible for differences in cardiac dysfunction and mortality risk.

Patients with a high CRF also had smaller left-sided C-IMT measurements compared to patients with moderate or low CRF. This may be because patients with a high CRF also had a better cardiometabolic profile, including high levels of self-reported physical activity. Exercise training can improve cardiovascular risk factor profiles in patients with CHD [39] and attenuate C-IMT and carotid plaque progression in as little as 6 to 12 months [40, 41]. It is however important to acknowledge that older age attenuated the relationship between C-IMT and CRF.

Conclusions

Identification of patients with low CRF and CHD appears to be an effective means of identifying those at highest

cardiometabolic risk, and increased risk of 5-year all-cause mortality. Exercise testing is widely applied in cardiological and CR environments. Longer-term, or higher intensity exercise-based CR programmes may help improve the cardiometabolic health of patients with CHD and low CRF.

Abbreviations

± : Standard deviation; 95% CI : 95% confidence Intervals; ANCOVA: Analysis of covariance; ANOVA: Analysis of variance; BMI: Body mass index; BP: Blood pressure; CABG: Coronary artery bypass graft; CCA: Common carotid artery; CHD: Coronary heart disease; CHF: Chronic heart failure; C-IMT: Carotid intima-media thickness; CPET: Cardiopulmonary exercise test; CR: Cardiac rehabilitation; CRF: Cardiorespiratory fitness; CV: Cardiovascular; CVD: Cardiovascular disease; DXA: Dual X-ray absorptiometry; ECG: Electrocardiogram; EDTA: Ethylenediaminetetraacetic acid; e-GFR: Estimated glomerular filtration rate; Hb: Haemoglobin; Hct : Haematocrit; HDL: High-density lipoproteins; HR: Heart rate; hs-CRP: High-sensitivity C-reactive protein; LDL: Low-density lipoproteins; LV: Left ventricular; LVEF: Left ventricular ejection fraction; METs: Metabolic equivalents; MI: Myocardial infarction; NHS: National Health Service; NT-proBNP: Terminal pro B type natriuretic peptide; VO_2/HR : Oxygen pulse; OUES: Oxygen uptake efficiency slope; PCI: Percutaneous coronary intervention; RPE: Rating of perceived exertion; SST: Serum separating tube; VAT: Ventilatory anaerobic threshold; VE/VCO_2 slope: Ventilatory efficiency with respect to CO_2 elimination; VO_2 : Oxygen uptake; $\text{VO}_{2\text{peak}}$: Peak oxygen uptake; η_p^2 : Partial eta squared

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Availability of Data and Materials

The dataset supporting the conclusions of this article are available via request to the corresponding author.

Authors' Contributions

SN contributed to study design, study management, data collection, data analysis, and manuscript preparation. CT, RP, AKB, and FN provided extensive support during data collection and manuscript preparation. TG contributed significantly to the study design, trial management, and manuscript preparation. ALC contributed to the study design and provided critical manuscript preparation as well as providing medical support to the study. SC contributed to the study design and data analysis, and reviewed multiple versions of the manuscript. LI contributed significantly to the study design and preparation, managed the study, and contributed to the manuscript preparation. All authors read and approved the final manuscript

Ethical Approval and Consent to Participate

Ethical approval for the study was given by the Humber Bridge NHS Research Ethics Committee-Yorkshire and the Humber (12/YH/0278). After giving verbal consent, patients were invited for assessment at Academic Cardiology, Castle Hill Hospital, Kingston-Upon-Hull, where written informed consent was obtained.

Competing Interests

Partial salary funding for SN was received by City Health Care Partnership CIC (Hull, United Kingdom). The authors declare that they have no competing interests.

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