High-intensity resistance training alone or combined with aerobic training improves strength, heart function and collagen in rats with heart failure

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Abstract: Background: Resistance training (RT) has been associated with positive responses in patients with cardiovascular disease, and when it is combined with continuous aerobic training (CAT), favorable adaptations appear to be even more pronounced. However, the effects of high-intensity RT alone or in combined with CAT in the case of heart failure (HF) is not completely elucidated. Methods: 28 male Wistar rats with HF (90 days old) were allocated to 4 groups: high-intensity RT (RT, n=7), CAT (CAT, n=7), RT and CAT (RT+CAT, n=7) and sedentary (Sed, n=7). Trained animals were subjected to a RT protocol in an adapted squat apparatus for rats (4 bouts, 6-8 reps, 90 s interval, 3×/week, 75% to 85% of one maximum repetition (1RM) for 8 weeks). The animals subjected to CAT performed it 3×/week during 50 min/session at 16 m/min. The animals of the combined exercise regimen performed both the RT and CAT exercise protocols. Results: The left ventricular end-diastolic pressure (LVEDP), collagen volume fraction and right ventricular hypertrophy were lower in RT, CAT and RT+CAT groups when compared to Sed group (P<0.05) for all outcomes. Regarding the inflammatory profile, only the CAT group showed greater IL-10 concentrations. Conclusion: We concluded that RT combined with CAT was able to improve the strength in animals with HF, which was associated to improvement in ventricular structure and function.

Keywords: Muscle strength, exercise, hemodynamic function, collagen, acute myocardial infarction

Introduction

Heart failure (HF) is related to a cardiac muscle modifies that lead to a systemic disorders, consequently yielding symptoms such as dysnea, fatigue and exercise intolerance [1]. Currently, muscle weakness has become a major target of research because the development of skeletal muscle strength is associated with survival of patients with HF [2, 3].

Changes in the extracellular matrix of cardiac collagen are connected with impairments in cardiac compliance, systolic and diastolic function and in the electrical behavior of the heart [4, 5]. Interestingly, physical exercise has produced favorable responses on interstitial collagen fractions in animal models [6, 7] and physical exercise has been described as an effective tool to implement cardiac reverse remodeling in the pathophysiology of HF [8].

Various exercise protocols have been employed as a strong non-pharmacological strategy in animal models of acute myocardial infarction (AMI) [6, 7, 9]. The use of resistance training (RT) in the experimental model with healthy rats has exhibited value as a tool for positive adaptations with respect to cardiac function, cardiovascular remodeling [10] and skeletal muscle [11].

The effects of aerobic and anaerobic physical training and its variables as type, frequency and intensity need to be elucidated [12]. Several studies using animal models of HF have employed the effects of continuous aerobic training (CAT) [7, 13]. However, the effects of high-intensity RT combined with aerobic training of moderate intensity are scarce. Therefore the aim of present study was to evaluate whether such responses may be associated with an improvement in hemodynamic function, inflam-
Resistance training and aerobic exercise in heart failure rats

Methods

This experimental study was based on ARRIVE (Animals in Research: Reporting In Vivo Experiments) guidelines [14].

Sample size, animals and ethical approval

Male Wistar rats (n=52, 220 to 270 g, 90 days of age) were obtained from the breeding animal unit of the Universidade Federal de Ciências da Saúde de Porto Alegre and were housed under standard conditions. This investigation followed the rules established by the guide for care and use of animals in experiments published by National Institute of Health (NIH publication no. 41-87, revised in 2011). All procedures were approved by the Ethics Committee in Research of Universidade Federal de Ciências da Saúde de Porto Alegre (protocol 199-13).

Acute myocardial infarction induction

The animals were weighed and anesthetized by inhalation with 2% isoflurane (Isoflurine, 100 ml-Cristália) in 100% oxygen [15] and were immediately artificially ventilated (Samway VR 15, 60 breaths/min). The ligature of coronary artery was performed to induce AMI as described previously [7].

Experimental design

Five weeks after AMI, 28 animals (with HF) were allocated into 4 groups: Sedentary (Sed), Resistance Training (RT), Continuous Aerobic Training (CAT) and Resistance Training combined with Continuous Aerobic Training (RT+CAT) (Figure 1).

Resistance training (RT)

Adaptation: The trained groups were subjected to a familiarization period with the adapted squat apparatus [16]. The animals were encouraged to perform 5 to 10 repetitions with 40% to 60% of body weight for one week [6].

1RM test: The training loads were determined based in percentage of 1RM test and adjusted every two weeks.

RT protocol: A neoprene vest attached to apparatus was used by animals to remain in the standard position on their lower limbs. An electric stimulus (4-5 mA, 1 s duration, with a 3 s interval between each repetition) was applied to the rats’ tails. The training protocol has been described elsewhere [6].

Continuous aerobic training (CAT)

Adaptation to treadmill: Five weeks after AMI induction surgery, the animals were encouraged to run (10 m/min; 0% inclination) for five minutes on the first and second day and ten minutes on the third, fourth and fifth days.

Exercise test tolerance/incremental load test: The animals were subjected to an exercise tolerance test (ETT) five weeks after AMI surgery and after eight weeks of training. The protocol employed consisted of a run on the treadmill with an inclination of 15° until exhaustion. The
Resistance training and aerobic exercise in heart failure rats

Protocol began with a speed of 5 m/min and there was an increment of 5 m/min every 3 minutes until exhaustion of the animal [17].

**CAT:** The run training protocol lasted for eight weeks, with 50 min/d; 3 times per week at 10 m/min, which was progressively increased up to 15 m/min (0% inclination).

**Protocol of combined physical training (RT+CAT)**

The Animals of the RT+CAT initially performed the RT and immediately at the end of RT, the animals were subjected to CAT.

**Evaluation of hemodynamic function**

The recordings were performed 48 hours after the end of the training protocol. The animals were anesthetized and a polyethylene catheter model P50 was introduced into the left ventricle (LV) via the right carotid artery. Blood pressure was recorded for 5 minutes immediately following the catheter being positioned within the LV to perform the registration of the ventricular pressure. This method has been described elsewhere [6].

**Blood sample and collection of tissue**

Blood collection was conducted with the tube already positioned in the left carotid artery and samples were taken in two 1.5 ml tubes containing 3.2% sodium citrate (1:9 v/v) and next centrifuged at 500xg for 10 min at 4°C. The plasma was stored at -22°C. Immediately after the hemodynamic assessment and blood collection, the animals were sacrificed by decapitation and the heart, lung and liver tissues were harvested [6].

**Cardiac hypertrophy, pulmonary and hepatic congestion**

Cardiac hypertrophy was determined by ratio to body weight [6, 7, 18]. To evaluate lung and liver congestion was utilized:

\[ \text{Congestion} \% H.O. = \frac{(\text{Wet weight} - \text{Dry weight})}{\text{Wet weight} \times 100} \]

**Determining the size of the infarcted area**

Left ventricles were filled with a latex pad and submerged in buffered formaldehyde for 10% over 24 h. The size of the infarcted area was determined by histological evaluation adapted from the method previously described by Martinez et al. [19].

**Analysis of interstitial collagen deposition in the heart muscle**

Cross cuts were performed on the LV and these fragments were fixed in 10% buffered formaldehyde for subsequent inclusion in paraffin [20, 21]. Histological sections of 6 microns were stained with picrosirius Red (PSR) and were employed for measurements of collagen volume fraction (CVF) present in LV. The fields selected were located far from the infarcted area and distant from the pericardial region as previously described [6]. The CVF was defined as the sum of all areas of interstitial collagen tissue (highlighted by polarization) divided by total tissue area [20].

**Determination of plasmatic levels of interleukins**

TNF-α and IL-10 plasma level were assessed through a Milliplex Map kit (RCYTO-80K, Millipore-Billerica, MA, USA) following the manufacturer’s recommendations.

**Statistical analysis**

Variables were presented as mean ± SD. After employing the normality test, ANOVA followed by a Tukey’s post hoc test to compare symmetrical variables between groups was used. A Two-way repeated measurement ANOVA followed by Tukey’s post hoc test was employed to compare the strength gain between groups at different experimental times and also to compare speed, distance and time in the ETT. Pearson’s correlation test was applied to examine the relationship among gain maximum strength and other variables. P values <0.05 were considered significant. GraphPad Prism version 6.01 (GraphPad Software, San Diego, CA, USA) for Windows. Sigmaplot version 12.0 (Systat Software Inc., San Jose, CA, USA) were utilized to statistical analysis.

**Results**

**Mortality, exclusion and adverse effects**

The mortality in AMI induction surgery was ~39% (20/52). The only animals with an infarct-
Resistance training and aerobic exercise in heart failure rats

The initial and final animal body weights as well other morphological variables were similar between groups in the pre- and post-training states. Sed-IC group showed greater cardiac hypertrophy and pulmonary congestion when compared to all training groups (Table 1).

**Morphological variables and pulmonary and hepatic congestion**

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**Hemodynamic evaluation**

RT group demonstrated lower LVEDP compared to Sed group. Higher value of -dP/dt_{max} was observed in RT+CAT compared to Sed group. The combined training protocol was also able to enhance cardiac relaxation as reflected by an increase of -dP/dt_{max}, mmHg/s (Table 2).

**Maximum strength gains: responses intra- and extra-groups**

Effects within groups: Figure 2 details the gain of absolute (A) and relative (B) force over time (1st to 4th 1RM). In terms of the 1st 1RM, there were no differences between groups in absolute and relative tests, suggesting that all groups started the protocol under similar conditions. The RT and RT+CAT groups presented gain maximum strength, in both, absolute and relative load in all tests compared to the initial test.
Resistance training and aerobic exercise in heart failure rats

Effects between groups: Figure 2A and 2B depict the absolute and relative strength gains between the groups. RT protocol alone or combined with CAT had an increase in their average values in the 2nd, 3rd and 4th 1RM compared to Sed and CAT groups.

Magnitude strength gain between the percentage differences of maximum load tests: Figure 2C and 2D details strength gains in the absolute and relative load in different moments of RM test for the RT and RT+CAT groups. These results suggest that during the protocol, there would be maximum strength gains, though on a smaller scale when compared to the effects of the initial two weeks of the RT protocol.

Pre-intervention no differences were observed between the groups, suggesting that all groups presented similar conditions at begin of study. CAT group showed an increase in speed, time and distance in post compared to pre-ETT. There were no differences between endurance-trained animals when compared to Sed and RT groups as well as CAT pre to post protocol.

Inflammatory profile

CAT group showed higher IL-10 levels compared to Sed group and there was no difference compared to RT and RT+CAT (Figure 4A and 4B). No differences were observed in both, TNF-α plasma level and TNF-α/IL-10 ratio between groups.

Interstitial collagen volume fraction

The RT isolated or combined with CAT groups demonstrates a lower collagen volume compared to Sed group (Figure 5).
Resistance training and aerobic exercise in heart failure rats

Correlations between central responses and maximum strength gain

Figure 6A-E, portray correlations between gain strength and the hemodynamic parameters and interstitial collagen fraction. There were significant correlation between LVEDP and the 4th 1RM (Figure 6A), LVEDP and the change in maximum strength gain (Figure 6B) and LVEDP and CVF (Figure 6C). In addition, there were significant correlations between CVF and the maximum strength gain as well as the change in maximum power gain (Figure 6D and 6E, respectively).

Discussion

To our knowledge, this is the first study to investigate the effects of 8 weeks of high-intensity RT (75% to 85% of 1RM) and CAT of moderate intensity (60% of the speed of ETT), alone or in combination, in HF rats. Based on our results, the trained groups showed improvement in the following variables: Strength gain, LVEDP, +dP/dtmax, -dP/dtmax, PC (% water), right ventricular hypertrophy, IL-10 and CVF. The maximum strength gain was an important outcome found in our study, because many studies have shown that muscle weakness is an independent risk factor for mortality in populations with HF [2, 3]. In this way, our study was able to show an increase in strength gain, which can be directly related to peripheral muscle preservation as a positive prognosis in HF syndrome.

Coronary artery ligation produces a marked ventricular dysfunction that is closely related to the infarct area size [6, 7] and therefore may be observed as a major impairment in hemodynamic function. In the present study, the infarcted areas were close to ~43%, which reflects directly a high LVEDP. Among the hemodynamic responses, we are able to highlight LVEDP, an important marker of ventricular dysfunction in HF [22].
Resistance training and aerobic exercise in heart failure rats

Figure 5. Collagen deposition analysis. Values are mean ± SD. A: Sedentary group with HF, Sed; B: Resistance training group, RT; C: Continuous aerobic training group, CAT; D: Resistance training combined with continuous aerobic training group, RT+CAT. E: Representative graph of the quantification of collagen volume fraction. *P<0.05 compared to Sed group.

Figure 6. Correlations between the left ventricle end-diastolic pressure (LVEDP) with maximum gain in strength and volume fraction of collagen. red circles (●), Sed; dark square (■), RT; dark triangle (▲), CAT; and blue triangle (▼), RT+CAT. A: LVEDP with 4th 1RM/BW; B: LVEDP with Δ maximum strength gains (%); C: LVEDP with fraction of collagen volume (%); D: Fraction of collagen volume with 4th 1RM/BW; E: Fraction of collagen volume with Δ maximum strength gains (%).
In our study, LVEDP [22] was lower in the RT (67.8%), CAT (48.7%) and RT+CAT (71.1%) groups compared to Sed group. We also saw that those animals that had an increase in muscular strength showed lower LVEDP values as demonstrated by the negative correlations in Figure 6A and 6B ($r=-0.62$ and $r=-0.73$, respectively). Previous studies in our laboratory with HF animals subjected to respiratory muscle training [23], swimming training [13] and CAT [7] showed similar responses in LVEDP. These improvements in cardiac function demonstrate a positive effect of exercise training.

The $+\Delta P/\Delta t_{\text{max}}$ and $-\Delta P/\Delta t_{\text{max}}$ exhibited favorable changes in the CAT group, while the group with combined exercise showed improvement solely in $-\Delta P/\Delta t_{\text{max}}$. Similar results have been reported in experimental studies of HF with different forms of interventions [7, 13]. In fact, the different exercise models like RT and CAT performed in our study leads to specific physiological adaptations on cardiac function. On the other hand, from a translational perspective, clinical studies have suggested there are controversial effects from the impact of exercise on systolic and diastolic function after acute myocardial infarction [12, 24, 25].

The exercise protocols used in the present study, were able to lower pulmonary congestion corroborating with other studies that employed RT or CAT as the primary intervention in rats with HF [6, 26]. Additionally, we observed lower values in the RVW/BW ratio in the RT+CAT group, which can be related to a cardiac remodeling associated to lower pulmonary congestion. These outcomes were also found in a previous study that utilized only the RT in rats with HF [6].

In our study, we observed an increase in plasma levels of IL-10 only in the CAT group. In fact, the moderate intensity of aerobic exercise is able to improve inflammatory profile [27]. Interestingly, the combined exercise group did not show improvement in this variable, possibly because the total volume and specific energy metabolism required in high intensity RT. These different characteristics of training may have interfered negatively to increased plasma levels of IL-10.

The modification of extracellular matrix collagen is part in a natural process of an adaptive response to increased mechanical load on the heart during exercise. However, the chronic overload arising from cardiac injury develops a negative response with respect to distensibility of the heart, valvular function, systolic and diastolic dysfunction and electrical transmission, ultimately leading to myocardial arrhythmia in HF [4, 5]. In the present study, the interstitial collagen accumulation of animals in the Sed group was higher when compared to groups subjected to physical training, demonstrating a possible reverse remodeling of the extracellular matrix, corroborating with the results of previous studies [26, 28]. These findings show that a sedentary behavior is directly related to a poor prognosis in HF condition. Here, a positive correlation was also observed between increased collagen accumulation and LVEDP ($r=0.64; P=0.0002$, Figure 6C). These results contributed to understand the relationship between hemodynamic injury and structural changes of the heart. Yet, it was observed in this study that an increase in muscle strength can influence the reduction of the accumulation of collagen as demonstrated by the correlation of Figure 6D and 6E ($r=-0.49$ and $r=-0.57; P=0.007$ for both, respectively).

There were some limitations in this study. First, the most robust biochemical analyzes in cardiac and peripheral muscle tissue could help to a better understand of possible balance between the molecular pathways of anabolism and catabolism induced by disease and exercise. Second, the absence of a load cell coupled to the squat apparatus for rats could help to increase the precision of maximum strength tests and control of the loads during the training protocol. This information could contribute to adjustments of intensity of training in the HF rats, especially in the RT training.

Conclusions

The RT protocol of high-intensity, performed alone or combined with CAT of moderate intensity, was able to increase the strength in animals with HF syndrome. This response was closely related to the improvement of ventricular structure and function. Nevertheless, the high-intensity RT protocol does not appear to offer benefits in terms of inflammatory markers (IL-10 and TNF-α). However, only the CAT protocol of moderate intensity was capable of exhibiting a positive influence on IL-10 plasma level.
We emphasize that the exercise protocols need to be better balanced in relation to the volume, intensity and frequency, since these variables can directly influence the functional and structural adaptations induced by physical training programs in HF.

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Disclosure of conflict of interest

None.

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References


Resistance training and aerobic exercise in heart failure rats


