



Review

Sex-Specific Impacts of Exercise on Cardiovascular Remodeling

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Abstract: Cardiovascular diseases (CVD) remain the leading cause of death in men and women. Biological sex plays a major role in cardiovascular physiology and pathological cardiovascular remodeling. Traditionally, pathological remodeling of cardiovascular system refers to the molecular, cellular, and morphological changes that result from insults, such as myocardial infarction or hypertension. Regular exercise training is known to induce physiological cardiovascular remodeling and beneficial functional adaptation of the cardiovascular apparatus. However, impact of exercise-induced cardiovascular remodeling and functional adaptation varies between males and females. This review aims to compare and contrast sex-specific manifestations of exercise-induced cardiovascular remodeling and functional adaptation. Specifically, we review (1) sex disparities in cardiovascular function, (2) influence of biological sex on exercise-induced cardiovascular remodeling and functional adaptation, and (3) sex-specific impacts of various types, intensities, and durations of exercise training on cardiovascular apparatus. The review highlights both animal and human studies in order to give an all-encompassing view of the exercise-induced sex differences in cardiovascular system and addresses the gaps in knowledge in the field.

Keywords: sex differences; exercise; cardiac remodeling; vascular remodeling; exercise-induced cardiovascular remodeling (EICR)



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1. Introduction

A topic of great interest over the past two decades has been the impact of exercise on the cardiovascular structure and function [1,2]. Exercise-induced cardiovascular remodeling (EICR) is often a result of adaptive, structural, functional, and molecular changes caused by exercise training. Although the term EICR is more recent, its study dates originated to over a century ago when researchers noted enlarged hearts in Nordic skiers and American university rowers [3,4]. Since then, several forms of EICR have been reported. Documented forms of EICR include chamber dilation, ventricular hypertrophy, increased coronary reserve [5], and enhanced diastolic filling [6]. Unlike the adverse outcomes of pathological cardiovascular remodeling, EICR often results in improved cardiovascular functions, including increased cardiac output and decreased resting heart rate. Furthermore, EICR has been shown to lower the risk of cardiovascular diseases (CVD), one of the leading causes of death worldwide [7], such as arrhythmias, heart failure, and ischemic conditions [8,9].

Another emerging area less well studied in cardiovascular remodeling is the impact of sex-specific exercise-induced alteration of cardiac and vascular structure [9]. It is known that cardiovascular structure and function differ between men and women. These differences in cardiovascular physiology are partly programmed by variation in male and female sex hormones [10]. For instance, stroke, hypertension, and atherosclerosis [11] are less common in healthy premenopausal women compared with age-matched men, but

these differences disappear in the postmenopausal years [12]. Furthermore, the decline in sex hormones with aging adversely remodels the cardiovascular system in a sex-specific manner [13].

The purpose of this review is to highlight sex disparities in the cardiovascular system and the sex-specific impacts of exercise in cardiovascular remodeling from both animal and human studies. We summarize potential sex-specific mechanisms underlying cardiovascular remodeling, shed light on the gaps in knowledge in the field, and propose future studies to advance the field.

2. Sex Disparity in Cardiac Structure and Function

It has long been acknowledged that sex differences exist in cardiovascular structure and function after adjusting for determining factors, such as height, weight, and body composition. In general, women have smaller and lighter hearts, and likewise, their coronary arteries are also smaller than their male counterparts [14,15]. Sex difference in the cardiac structure becomes apparent at puberty, coinciding with the rise in sex hormones. In the pubertal age group, left ventricular (LV) mass is greater in boys than that in girls and chamber dimension, wall thickness, and myocyte hypertrophy likely contribute to the higher LV mass in males beyond puberty [16,17].

As age advances, the male heart loses on an average, one gram of cardiomyocytes daily, while the adequate heart mass is maintained. However, adult female hearts exhibit more resistance to cardiac cell loss and hypertrophy compared to their male counterparts [17–19]. Female sex hormones in general and estrogen in particular are thought to contribute to the prevention of cardiac hypertrophy in females. Several studies have demonstrated that estrogen attenuates cardiomyocyte apoptosis and modulates both physiological and pathological LV hypertrophy [20–24]. On the contrary, the loss of cardiomyocytes in males may be secondary to elevated levels of testosterone and/or epinephrine. These hormones have been associated with apoptosis and fibrosis [25–27].

Similarly, there are sex differences in cardiac function. For instance, the diastolic function is greater in premenopausal women compared to age-matched men [17,28], but this difference disappears in the postmenopausal years. Though the diastolic function decreases in both sexes with aging, reduction in systolic function occurs mainly in men [17,19]. Furthermore, studies show that the LV dimensions and functions also vary among men and women. Women have smaller LV chambers and reduced stroke volume (SV) compared to men [29]. Therefore, women rely on a higher heart rate compared to men to maintain similar cardiac output.

Sex differences are also seen in cardiac metabolism. Cardiac function is dependent on substrate oxidation (fatty acid, glucose, lactate, ketone, triglyceride, and glycogen) for the production of ATP [30]. Sex hormones may play a role in energy metabolism in the myocardium. For instance, a study performed in twenty-five young, healthy adults showed that women's hearts use less glucose and more oxygen compared to the age-matched men [30]. Moreover, estrogen is known to increase endothelial NO synthase (eNOS), and its upregulation causes a decrease in translocation of glucose transporter (GLUT)-4 to the cell surface, thus inhibiting the glucose uptake and utilization in cardiomyocytes [11,31]. From these observations, it can be postulated that female hearts depend more on fatty-acid oxidation for myocardial energetics. Successively, they require more oxygen consumption, necessary for higher energy production, to meet the higher cardiac function compared to males [31]. Therefore, the sex-specific cardioprotection seen in females may only be limited to a well-perfused and oxygenated, healthy heart. In contrast, during cardiac stress and disease, the female myocardium may be less likely to adapt to make a shift in substrate metabolism.

3. Sex Disparity in Vascular Wall Structure and Function

There are several studies that show the protective role of estrogen in the vasculature, with fewer studies focusing on the impact of androgens on vasculature.

Male sex steroids have been linked to the enhancement of large artery stiffness [32]. During their reproductive years, female arteries exhibit lower arterial stiffness and prevalence of hypertension compared to age-matched males [33]. However, menopausal women exhibit stiffer large arteries than their male counterparts [32,33], manifesting in elevated prevalence of hypertension [34]. Berry et al. showed that elderly hypertensive women have stiffer large arteries, greater central wave reflection, and higher pulse pressure than elderly men [35]. Furthermore, in a longitudinal multiethnic cohort, Stern et al. reported that carotid arterial stiffening associated with aging could lead to an increase in systolic blood pressure in both sexes [36]; however, race and ethnicity (in men) and level of education (in women) could be further contributed to the differences between the sexes.

Many of estrogen's beneficial vascular effects are related to modifying the functional state of the endothelium [37]. A number of studies indicate that estrogen promotes the release of endothelium-derived relaxing factors that confer resistance against atherosclerotic events. It was previously reported that the administration of estrogen to ovariectomized rats improves vascular function through its effects on vasodilators, such as nitric oxide (NO) [38–40]. These findings have been confirmed in both human [20,41,42] and animal [43] studies.

The sex differences in vascular tone and hypertension may be also related to differences in the production of or sensitivity to endothelium-derived contracting factors, such as endothelin-1 (ET-1). The lower blood pressure in female compared with male spontaneously hypertensive rats (SHR) has been attributed to the decreased level of ET-1 in females [44]. Several studies have reported that the contraction to phenylephrine, an alpha adrenergic agonist, is greater in the aorta of the male than in that of the female rats [45–48].

Overall, there are sex differences in cardiovascular structure, function, and metabolism. Furthermore, sex-specific differences exist in CVD. Exercise has been shown to have beneficial effects on the cardiovascular system [8,49]. Understanding of sex differences in cardiovascular system provides important mechanistic information for sex-specific impacts of exercise in healthy and pathological conditions. Next, we will highlight the sex-specific impacts of exercise with relation to various types of exercise regimens on cardiovascular remodeling.

4. Exercise-Induced Cardiovascular Remodeling and Functional Adaptation

Various types and intensities of exercise uniquely determine the outcomes on cardiovascular remodeling, so it is important to define the different categories of exercise in order to understand its impact on the cardiovascular system [9,50]. Exercise can be defined as a form of physical activity involving planned, structured, and repetitive movements, with the intention of maintaining or improving physical fitness [51]. Often, both intensity and duration are calculated based on the body mass index and age of men and women. Exercise intensity may be divided into three categories (low, moderate, and high intensity, see Table 1) based on the metabolic equivalent of task value (MET). These categories are as follows: low (<3 METs), moderate (3–5.9 METs), and high/vigorous (≥ 6 METs) [52]. Exercise intensity can also be quantified by the maximum oxygen consumption (VO_{2max}) or the maximum heart rate (HR_{max}) during the exercise. For low to moderate exercise, this would be defined as approximately 45–70% of VO_{2max} or 55–74% of HR_{max} respectively [53,54]. For high-intensity exercise, this would be defined as greater than 70% of VO_{2max} or greater than 90% of HR_{max} [53–55]. Of interest to this review, women exhibit a (5–15%) lower VO_{2max} compared with men when controlled for age and weight or activity [56]. A lower cardiac output is shown to contribute to a lower VO_{2max} in women in some studies [57,58]. Some pre-clinical and clinical studies have also characterized exercise training by endurance and strength training. It has been shown that various forms of exercise induce different types of EICR [59]. In this review, we will focus on exercise regimens including aerobic exercise (low–moderate and high-intensity) and combined exercise (aerobic and resistance training) (see Table 2).

Table 1. Commonly accepted definitions of the different intensity levels of aerobic exercise.

Exercise Intensity	MET *	VO _{2max} *	HR _{max} *
Low	<3	<45%	<55%
Moderate	3–5.9	45–70%	55–74%
High	≥6	≥70%	≥90%

MET, metabolic equivalent of task; VO_{2max}, maximum oxygen consumption; HR_{max}, maximum heart rate.
* Approximate ranges.

Table 2. Definitions of the different types of exercise.

Types of Exercise	
Aerobic	Activity that increases the capacity of the cardiorespiratory system by increasing oxygen supply and improving the oxygen utilization in muscles. Can be categorized further by intensity, such as low, moderate, and high.
Resistance	Exercise that includes the use of a load, machinery, or your own body weight to increase muscle strength and endurance. Strength and endurance training are subtypes of resistance training.
Combined	An exercise routine that incorporates a combination of aerobic and resistance exercise.

In animal studies, unlike humans, exercise-training conditions are precisely defined and include forced treadmill training protocols, swimming, and cage wheel running [60]. Several animal studies have shown sex-specific differences in physiological hypertrophy with exercise. For instance, female rodents show greater physiological hypertrophy compared to male rodents with both types of swim and treadmill training [60–63]. Interestingly, female rodents can run faster and for greater duration compared to males when controlled for age and strain [63]. Furthermore, key metabolic pathways, including phosphoinositide-3 kinase (PI₃K)/Protein Kinase B (AKT) pathway and calcium-calmodulin dependent kinase (CaMK) pathways, are impacted by exercise and associated with physiological hypertrophy in animal studies [8,49]. Exercised female groups also exhibit an increased phosphorylation of a negative inhibitor of physiological hypertrophy, Glycogen Synthase Kinase-3-beta (GSK-3-beta), an upstream mediator of AKT phosphorylation, that led to increased hypertrophic growth in the hearts of this group [63]. Furthermore, exercise selectively upregulates CaMK pathway in female rodents compared to exercised male rodents [63]. In contrast, sedentary male rodents exhibit selective upregulation of CaMK pathway compared to sedentary females [63].

Renin–angiotensin system (RAS) also plays an important role in the progression of cardiac remodeling. A decrease in the angiotensin-converting enzyme (ACE)/angiotensin (Ang) II/Ang II type 1 receptor (AT1) axis of RAS provides protection from pathological cardiac hypertrophy and subsequent heart failure [64,65]. Activation of AT1 receptor by Ang II has been shown to increase collagen and myocyte hypertrophy [64]. However, activation of angiotensin-converting enzyme 2 (ACE2) leads to the formation of Angiotensin (1–7)/Mas, which exhibits vasodilatory, anti-proliferative, and anti-trophic effects [66–68]. One study showed that six weeks of swimming training in FVB/N mice lacking Mas induced cardiac hypertrophy, which was associated to an increase in collagen I and III mRNA expression [66]. The investigators suggested that the increase in collagen attributed to an inversion of the balance between Ang II and Angiotensin (1–7) actions in the heart of Mas-knockout mice, favoring a stronger and unopposed influence of Ang II. Their data indicate that Angiotensin (1–7)/Mas axis is an important counter-regulatory mechanism in physical training-mediated cardiac adaptations [66].

In human studies, concentric thickening of LV is observed in strength training, whereas eccentric increase in cavity size of LV is more pronounced in endurance training [59,69]. There are structural and functional differences between the cardiovascular systems of young, trained athletes and the untrained ones. The LV cavity dimension and maximal wall thickness are greater in athletic females than the nonathletic females [70]. Moreover, the remodeling is different in males and females. The LV cavity dimension and wall thickness

are smaller in female athletes compared to the male athletes of the same age, race, and sport disciplines [70]. Furthermore, women tend to develop eccentric hypertrophy, while concentric hypertrophy may be a normal finding in male athletes [71]. Women also exhibit improved endurance capacity and have increased catecholamine-induced fatty acid uptake and oxidation with exercise training compared to males [72,73]. Sex differences are also seen in SV with aging during submaximal and maximal exercise; older women have a lower SV and a smaller increase in SV from rest compared to aging men. This difference in SV is thought to in part be secondary to differences in LV remodeling with exercise and differences in lean body mass between the sexes [56]. In contrast, there is no difference in SV, cardiac index, LV end diastolic, and systolic volume indexes in younger (<40 years old) men and women [56].

Interestingly, exercise has variable impact on different heart chambers. For instance, intensive training-induced hemodynamic changes have been found to cause atrial enlargement in competitive athletes. A meta-analysis conducted by Iskandar et al. demonstrated that elite athletes have larger left atrium (LA) diameter and volume compared to control subjects [74]. The atrial enlargement, which is dynamic and reversible, is an adaptive mechanism in response to the increased training-induced volume overload, and it may vary by sex. In a study consisting of young elite rowers (46.5% women), LA enlargement was found more frequently in men than in women [75]. Similar to the LA, greater right atrium (RA) dimensions were observed in elite athletes compared to sedentary control subjects [76]. In addition, sex differences in EICR were also found in a study conducted in highly trained university athletes. While the male athletes showed LA and LV remodeling, right ventricular (RV) remodeling was significantly more common in female athletes [77]. Overall, the impact of exercise on sex-specific cardiac remodeling is greater in untrained men and women than in highly trained athletes [56].

Beneficial effects of exercise are also noted in the vascular endothelium [78]. For instance, the levels of endothelial-derived ET-1 and NO are impacted by exercise [79,80]. Exercise-induced increased cardiac output and intermittent increase in laminar shear stress contribute to vascular endothelial remodeling by activation of the PI₃K/AKT pathway, leading to eNOS phosphorylation and subsequent increase in NO production [80]. NO-mediated endothelium-dependent vasorelaxation is also improved with exercise training [78]. NO is a key mediator of angiogenesis [80,81]. Up-regulation of vascular endothelial growth factor (VEGF) during exercise is required for angiogenesis associated with exercise [80]. However, one study in the literature reported both the basal levels of VEGF and response to exercise in the patients with cardiac failure was not different from the controls [82]. In a study of healthy male subjects, strength training led to an increase in ET-1, leading to increase in aortic pulse wave velocity (PWV), an established indicator of vascular stiffness. On the other hand, the same study showed that endurance training down regulated ET-1 and thereby reduced arterial stiffness [79]. Vascular endothelial dysfunction is a hallmark of the vascular disease; it is defined as a reduced endothelium-dependent vasodilation to vasodilators, such as acetylcholine (ACh) or flow-mediated dilation (FMD). Thus, the responses of vessels to ACh or FMD are used as a reproducible parameter to investigate endothelial function following exercise.

In a study of aging adults, twenty-four weeks of exercise led to improvement in vascular FMD and function in women but not in men [83]. In contrast, eight weeks of exercise led to the improvement of FMD in aging men but not women [84], suggesting that the duration of exercise may play a role in its sex-specific outcome. Another study in the literature demonstrated that exercise led to an increase in shear stress-mediated vasorelaxation in the brachial artery; however, this study did not include female subjects [85]. Overall, sex-specific impacts of exercise have been studied in the vasculature; however, data are sparse. Interestingly, beneficial roles of estrogen in combination with exercise are reported in postmenopausal women. In a study by Moreau et al., endurance training for twelve weeks in conjunction with estrogen treatment increased FMD in brachial arteries of postmenopausal women [86].

Beneficial impacts of exercise on cardiovascular structure and function are also seen in CVD, such as ischemia-reperfusion injury (IRI) and chronic heart failure (CHF). Endurance exercise has been reported to reduce oxidative stress and structural damage in IRI, thereby preventing myocardial dysfunction in animal studies [87–92]. However, the role of biological sex (if any) on the effects of exercise on myocardial oxidative stress is not clear. This is simply because the majority of studies did not include both sexes. Endurance training is also shown to reverse LV remodeling, with modest improvement in ejection fraction (EF) and LV end diastolic volume in CHF patients [93,94]. These studies were, however, performed only in males or the sex of the enrolled patients was not noted.

Animal studies provide further mechanistic insights into attenuation of adverse remodeling associated with heart failure. In a rodent model of dilated cardiomyopathy, swim training activated PI₃K and improved survival. However, this investigation was performed only in males, and females were not studied [95]. Exercise has also been reported to upregulate sarco/endoplasmic reticulum Ca²⁺-ATPase (SERCA) pump in pathological hypertrophy, thus improving calcium handling and contraction in animal models [96,97]. Moreover, it is shown that CHF-induced increased sympathetic activity is reduced with exercise. For instance, eight weeks of exercise training reduced norepinephrine secretions by 16% [98]. Braith et al. demonstrated that endurance exercise reduced neuroendocrine activity by reducing the neuroendocrine hormones, including Ang II, vasopressin, and atrial natriuretic peptide; sex differences were not addressed [99]. Cardiac-failure-related increase in collagen deposition was also reduced in exercised animals; only males were studied [100].

Exercise has also shown to have beneficial effects on vascular diseases. Hambrecht et al. noted that four weeks of endurance exercise led to increase in phosphorylation of eNOS in the vessels of patients with coronary artery disease (CAD), thereby improving vasodilatory capacity [101]. Similarly, four weeks of exercise reduced Ang II in the vessels of male patients with CAD leading to reduced vasoconstriction; females were not studied [102].

5. Detrimental Impacts of Exercise

Despite the benefits of exercise on cardiovascular remodeling, there are also concerns that athletes are predisposed to the pathological cardiac remodeling, cardiovascular dysfunction, and arrhythmias.

5.1. Detrimental Impacts of Exercise on Cardiac Structure and Function

A meta-analysis reported that prolonged endurance exercise reduced RV function without causing any alteration to LV function in healthy individuals over 18 years of age [103]. A greater pulmonary artery pressure during exercise increases RV end-systolic wall stress, causing greater reduction in RV function [103]. Because exercise can impose a disproportionate physiological load on the RV compared to the LV, intense and prolonged exercise may result in long-term cardiac fatigue and remodeling [104–106]. A previous study suggests that some athletes suffer from serious arrhythmias emerging from pathological RV remodeling caused by chronic endurance exercise [107]. In fact, arrhythmogenic RV cardiomyopathy (ARVC) is an example of pathological remodeling and considered as one of the leading causes of sudden death in athletes [69,108–110]. However, since intense exercise may also lead to physiological adaptations of the heart, it could be difficult to differentiate between physiological RV remodeling and early-stage ARVC [69,110]. Cardiac magnetic resonance (CMR) has been utilized as a gold standard to assess RV structure and function and may help differentiate pathological remodeling from physiological remodeling [111].

5.2. Detrimental Impacts of Exercise on Cardiac Electrical Conductivity

Studies have shown an increased risk for developing atrial fibrillation (AF) in endurance athletes than in non-athletes [111,112]. Although data on the cardiac impact of endurance exercise in women athletes are scarce, studies have proposed that women are

more protected from developing AF [112]. This sexual dimorphism could be explained by the difference in electrophysiological changes that occur in men and women during exercise. For instance, Wilhelm and co-workers observed significantly longer signal-averaged electrocardiogram (ECG) P-wave duration and increased incidence of AF in male cyclists compared to women cyclists [113]. The link between longer P-wave duration and increased risk of AF is already well known [114]. Animal studies have also corroborated with increased susceptibility to AF with chronic endurance exercise [115]. Moreover, underlying cardiac conductivity defects, like prolonged QTc and Brugada syndrome, can also lead to cardiac arrhythmias [116–120] and may have the potential to cause detrimental cardiac outcome with aerobic activities. It is also important to note that there may be sex differences in arrhythmogenic potentials secondary to underlying conditions, such as catecholaminergic polymorphic ventricular tachycardia (CPVT) associated with cardiac ryanodine receptor 2 (*RyR2*) mutations and other cardiac pathologies [116–125].

5.3. Detrimental Impacts of Exercise on Vascular Structure and Function

Although there have been studies that showcase the positive outcomes of exercise on vasculature, there are also reports on detrimental effects of high-intensity exercise [126]. For example, high-intensity exercise has been shown to decrease FMD in male participants [127,128]. However, there are also reports of increased FMD with high-intensity exercise in both sexes [56,129]. Overall, studies have shown that the alteration in FMD is dependent on exercise intensity [128,130]. In addition, there are reports on the exercise-induced increase in markers of endothelial dysfunction including, Von Willebrand factor, and thrombomodulin (in males) and microRNA-126 (in both sexes) [131,132]. Furthermore, a study by Boos et al. found evidence of exercise-induced vascular damage with an increase in circulating endothelial cells in men and women with CAD [133]. Contrasting effects of exercise on vascular function are further summarized in reports by Sapp and Hagberg, 2018, and Adams, 2018 [126,134]. Additional studies are needed to document the impact of exercise intensity on detrimental vascular outcomes.

6. Sex-Specific Impact of Low–Moderate Intensity Exercise on Cardiovascular Remodeling

Low to moderate-intensity exercise includes any activity that uses equal or less than 5.9 METs. In humans, exercises such as walking (up to 6 km/h), dancing (3.3 METs), or stationary cycling with light effort (at 100 Watts is equal to 5.5 METs) are considered as low–moderate intensity exercises [52,135]. However, in animal models, the measure of intensity is frequently based on minutes per day of activity. For instance, in rodents, ≤ 90 min/day of swimming or voluntary cage wheel running are considered low to moderate exercise [136]. Table 3 contains details of the low–moderate intensity training regimens and the studies discussed below.

6.1. Sex Disparity in Cardiac Structure and Function with Low–Moderate Exercise

In this section, we will discuss both animal and human studies addressing sex disparities in cardiac structural and functional remodeling with low–moderate intensity exercise.

Animal studies have shown that moderate-intensity training (MIT) induces physiological cardiac hypertrophy [54,137]. Asif et al., in a study only examining male rats, showed that MIT led to an increase in cardiac mass. This study also showed an age-specific impact on EICR by demonstrating that exercise led to increased LV mass in both adult and adolescent rats, but an increase in the number of cardiomyocytes was only seen in adolescents [137]. Apart from the induction of physiological hypertrophy, MIT can also reduce age-induced pathological cardiac hypertrophy. A study looking at only aged male rats exhibited a reduction in their pathological cardiac hypertrophy when they underwent MIT [138]. Furthermore, the same study showed that MIT training led to reduction in cardiac inflammation and cardiac fibrosis in aged male rats [138].

Molecular mechanisms underlying the beneficial effects of MIT on physiological cardiac remodeling have been also described. Dworatzek et al. demonstrated that mice

undergoing MIT exhibited the activation of PI₃K/AKT signaling pathway in the LV, thereby leading to cardiac hypertrophy. However, when compared to males, female mice exhibited greater cardiac hypertrophy following MIT [139]. Furthermore, in the same study, estrogen beta receptor knockout female mice had comparable cardiac structural change to the males, suggesting that sex differences observed in EICR are, in part, mediated via estrogen receptor signaling [139]. Beneficial effects of low–moderate exercise on cardiac function can be also mediated by other molecular mechanisms. For instance, rats undergoing MIT exhibited a decrease in collagen and an increase in citrate synthase levels in the LV [140]. Citrate synthase is necessary for aerobic capacity and mitochondrial mass, while increased collagen levels are associated with increased myocardial stiffness and heart failure [140,141]. Similarly, oxidative stress is known to contribute to pathological cardiac remodeling, and it is known that exercise training reduces oxidative stress in the heart [8].

Low to moderate-intensity exercise is shown to have several beneficial effects on cardiac function in animal studies. For instance, Hafstad et al. demonstrated that diet-induced obesity resulted in diastolic and systolic dysfunction in mice, and that could be prevented with MIT. However, females were not included in this study [142]. In another study, MIT resulted in decreased end systolic volume, increased SV, increased EF, and decreased LV pressure in male rats [140].

The effects of low–moderate intensity exercise on the cardiovascular system are not limited to animal models, as similar effects have also been demonstrated in human studies. In Dawes et al.'s study, moderate intensity, defined as exercise between three to five hours per week (assessed by a questionnaire), was associated with increased LV mass and increased LV and RV volume in healthy adult men and women; however, sex differences were not addressed, as patient data were grouped [143]. Another study found that in a community setting, intentional exercise, defined as the sum of activities that were consciously done for exercising, such as sports/dancing, conditioning activities, and walking regardless of the intensity level, was associated with increased LV mass, increased SV, and increased end diastolic volume in both men and women. Furthermore, the rate of increase in LV mass, SV, and end diastolic volume were more pronounced during lower intensity training in both sexes; however, the magnitude of changes was smaller in women than in men [9]. The underlying reason(s) for the sex differences in the beneficial impact of lower-intensity training in this study were unknown. However, the investigators made several speculations, including that the increased ventricular mass in men may be attributed to the larger hearts in males (despite accounting for baseline heart mass). Moreover, they suggested that the augmented changes in ventricular mass in men may have also involved participating in higher intensity of exercise compared to females [9].

6.2. Sex Disparity in Vascular Structure and Function with Low–Moderate Exercise

Similar to the beneficial impacts of low–moderate exercise on the heart, the beneficial impacts of exercise have been shown on vasculature. For instance, in a study only looking at male rats, Patora et al. demonstrated that MIT resulted in hypertrophy and morphologic changes in aortic smooth muscle cells as well as an increase in the thickness of aortic elastic fibers. This increase in elastic fiber thickness was thought to lead to an increase in vascular distensibility, making the vessels better at handling mechanical stress [144].

Functional changes in the vasculature have also been observed with low–moderate intensity exercise. There has been increasing evidence suggesting a link between exercise and vascular ion channels [145–147]. Zhang et al. noted that aerobic exercise lowered systemic blood pressure and normalized hypertension-associated large-conductance calcium-activated potassium channels (BK_{Ca}) upregulation to normotensive control levels in male SHR, and these effects were more pronounced in the moderate-intensity group than in the low-intensity group [148].

Table 3. The described studies associated with the low–moderate intensity training regimens.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanisms
Asif et al. [137]	Male Wistar Kyoto rats	Running: treadmill Duration: up to 1 h/day, 5 days/week for 4 weeks	↑LV diameter	N/A	↓Cardiac microRNA-208b in LV (authors believe this change to be insignificant)
Liao et al. [54]	Aged male Sprague-Dawley rats	Swimming Duration: gradually increased from 20 to 60 min/day, 5 days/week for 12 weeks	↓Pathological cardiac hypertrophy	N/A	Down regulation of ERK1/2/JNK and NFATc3
Dworatzek et al. [139]	Male and female C57BL/6J mice	Running: voluntary cage wheel Duration: 8 weeks	↑LV mass	Greater cardiac hypertrophy in females compared to males	Activation of PI ₃ K/AKT signaling pathway by upregulating AKT/mTOR signaling leading to cardiac hypertrophy
Verboven et al. [140]	Male Sprague-Dawley rats	Running: treadmill Duration: 1 h/day, 5 days/week for 13 weeks	↓End systolic volume, ↑SV, ↑EF, and ↓LV pressure	N/A	N/A
Hafstad et al. [142]	Male C57BL/6J mice	Running: treadmill Duration: ~120 min/day, 5 days/week for 10 weeks	Prevention of diet induced diastolic and systolic dysfunction	N/A	N/A
Potora et al. [144]	Male Wistar rats	Swimming Duration: 15 min/day for 14 days	Aortic smooth muscle cells hypertrophy and morphological changes, ↑thickness of elastic fibers	N/A	N/A
Zhang et al. [148]	Male SHR	Running: treadmill Duration: 60 min/day, 5 days/week for 8 weeks	↓Systemic BP	N/A	Correcting the hypertension-associated BK _{Ca} channel remodeling and suppressing the pathological adaptations of BK _{Ca} channels that result from high BP

Table 3. Cont.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanisms
Dawes et al. [143]	Males and females	MIT as defined by the Copenhagen City Heart Study Leisure Time Physical Activity Questionnaire	↑LV mass and ↑LV and RV volume in males and females	Data were not analyzed for sex differences	N/A
Turkbey et al. [9]	Males and females	MIT as defined by the MESA Typical Week Physical Activity Survey	↑LV mass, ↑SV, and ↑end diastolic volume in both sexes	Males showed a greater increase in LV mass, SV, and end diastolic volume as the levels of physical activity increased when compared to females	N/A
Shenouda et al. [149]	Healthy males	Stationary cycling Duration: 45 min/day for 12 weeks	↑Brachial artery FMD but no change in PWV	N/A	N/A
Sawyer et al. [150]	Healthy males	Stationary cycling Duration: 40 min/day, 3 days/week for 8 weeks	↑Brachial artery diameter but no significant change in FMD	N/A	N/A
Rakobowchuk et al. [151]	Males and females	Stationary cycling Duration: 40–60 min/day, 5 days/week for 6 weeks	↑Relative FMD and improved distensibility in popliteal artery	No sex differences were seen	N/A
Goto et al. [152]	Healthy males	Stationary cycling Duration: 30 min/day, 5–7 days/week for 12 weeks	↑Endothelium-dependent vasodilation	N/A	↑Production of NO
Sugawara et al. [153]	Post-menopausal females	Stationary cycling Duration: 3–5 days/week for 12 weeks	↑Arterial compliance and ↓ LDL	N/A	N/A

↓decrease, ↑increase, left ventricular (LV), stroke volume (SV), ejection fraction (EF), spontaneously hypertensive rats (SHR), blood pressure (BP), large-conductance Ca²⁺-activated K⁺ channel (BK_{Ca}), moderate-intensity training (MIT), right ventricular (RV), flow-mediated dilation (FMD), pulse wave velocity (PWV), nitric oxide (NO), low-density lipoprotein (LDL).

It has also been shown that low–moderate intensity exercise has beneficial effects on the peripheral vasculature in humans. One study in males demonstrated that moderate intensity exercise in the form of stationary cycling increases resting brachial artery diameter by 8%. Of note, brachial artery diameter is a measure of peripheral vessel function [149]. However, MIT did not result in changes in any of the other recorded cardiovascular measures, such as arterial stiffness, resting blood pressure, and heart rate; females were not studied, though [149]. The increase in brachial artery diameter noted with MIT is suggestive of structural remodeling that may be an adaptation to meet the blood-flow demands of exercise [149]. A similar study by Sawyer et al. also found that MIT resulted in increased brachial artery diameter. This study used both male and female subjects, but the investigators did not analyze them in separate groups; thus, no sex differences were elicited [150]. Another study that looked at the effects of high-volume endurance training (at 65% of VO_{2max}) showed that popliteal artery distensibility and popliteal endothelial function were both improved [151]. However, no sex differences were observed in either popliteal artery distensibility or popliteal endothelial function with this mode of exercise [151]. In another study by Goto et al., MIT resulted in increased forearm blood flow in response to ACh through the increased production of NO. However, this study looked only at men [152]. Similarly, Sugawara and coworkers demonstrated that in post-menopausal women, both low and moderate-intensity exercise resulted in increased carotid arterial compliance. However, resting heart rate, blood pressure, and pulse pressure were not significantly changed in either low or moderate-intensity groups [153]. Furthermore, Sugawara and coworkers reported that both low and moderate-intensity exercise decreased low-density lipoprotein (LDL) but not high-density lipoprotein (HDL) cholesterol in post-menopausal women [153].

6.3. Limitations

The data detailing the impacts of low-moderate exercise on cardiovascular remodeling are limited. In particular, fewer studies address the effects of MIT in humans as compared to small animal models. Therefore, there is a need to translate some of the preclinical findings to humans and design prospective studies to address this gap. Another challenge in evaluating the effects of exercise on the cardiovascular system is the failure to uniformly define exercise intensity. In this review of the literature, we noted that some authors failed to report the VO_{2max} or METs used during training. Preclinical and clinical research should consider utilizing METs and VO_{2max} to uniformly and accurately define low-moderate intensity exercise to assess its impact on cardiovascular remodeling. Furthermore, there is also need for more prospective studies on low-moderate exercise that include both male and female subjects and assess sex-specific impact of exercise on EICR. The sparse data currently available suggest that there are sex differences with low-moderate intensity exercise in cardiac remodeling; however, we are not able to ascertain similar differences in vascular remodeling [9,139].

7. Sex-Specific Impact of High-Intensity Training (HIT) on Cardiovascular Remodeling

HIT or vigorous exercise involves physical activity that requires ≥ 6 METs or greater than 70% of VO_{2max} or over 90% of maximal heart rate [52–55]. World Health Organization's definition of vigorous exercise includes running, fast cycling, fast swimming, sports, or weightlifting that require greater effort and increased breathing and heart rate [154]. According to the U.S. Department of Health and Human Services 2018, 75–150 min/week of HIT is sufficient for substantial health benefit outcomes [155]. HIT has recently gained popularity in the form of high-intensity interval or intermittent training (HIIT). HIIT is defined as an exercise that combines relatively short to long bursts of intense exercise with periods of rest or lower intensity exercise [156,157]. Though the workout period is relatively short, studies showed that HIIT produces health benefits similar to or greater than MIT [157–160]. HIT is a well-known therapeutic intervention that results in physiological cardiovascular remodeling and has proven therapeutic benefits in pathological cardio-

vascular remodeling in both animal models and human subjects [129,140,156,161–163]. However, controversies exist, as both beneficial and detrimental effects on cardiovascular system have been reported with HIT. Table 4 contains details of the HIT regimens and the studies discussed below.

7.1. Sex Disparity in Cardiac Structure and Function with HIT

Cardiac structural and functional benefits have been noted in the literature with both HIT and HIIT. Preclinical studies are suggestive of the beneficial impacts of HIT on physiological cardiac remodeling in a healthy state. One study in the literature found that high-intensity swimming resulted in increased phosphorylation of AKT in the myocardium in both male and female rats but to a greater degree in females, thus leading to more pronounced LV hypertrophy in females [164]. HIT also increased SV and improved contractility and stroke work in both sexes; however, improvement in diastolic function was only seen in male rats [164]. Similarly, in a study of healthy male rats (females were not included), high-intensity treadmill running induced physiological LV hypertrophy. Moreover, improved cardiac performance (including enhanced EF, cardiac output, and volume), reduced myocardial collagen content, and increased cardiac capillary density were observed [140]. In addition, de Oliveira et al. demonstrated that, in a study of only male mice overfed a diet high in fat or fructose, HIIT reduced LV mass and LV wall thickness. These beneficial effects of HIIT were attributed to elevation in the components of the cardiac RAS, ACE2/Angiotensin (1–7)/Mas receptor [156]. As previously stated, activation of Angiotensin (1–7) in the myocardium is associated with vasodilation and antifibrotic, anti-hypertrophic, and antiarrhythmic actions [165,166]. RAS further modulates ACE2/Angiotensin (1–7)/Mas receptor axis, thereby exerting an anti-inflammatory effect in the myocardium, leading to beneficial impact on cardiac remodeling [167,168].

HIT has also been reported to have favorable effects in the disease states. Exclusively in a male rat model of pulmonary hypertension, high-intensity treadmill training enhanced RV apelin (a potent vasodilator) expression [169], leading to decreased RV systolic pressure, RV hypertrophy, fibrosis, and improved cardiac output [161]. HIIT training was also reported to have cardioprotective effects in male rats with IRI [170]. Rahimi and coworkers showed that the infarct size in the exercised male groups reduced by 50% and 35% after one and seven days post exercise, respectively, compared to the sedentary group [170]; females were not included. However, the beneficial effects of HIT on ischemic heart were lost after fourteen days following detraining. These investigators, therefore, proposed that the beneficial impacts of short-term HIIT may persist for a short period following exercise [170]. Of note, detraining is a state of decline in exercise-induced physiological conditioning due to insufficient training or pause in training.

In a study performed in healthy, middle-aged male individuals, HIIT induced beneficial RV hypertrophy, improved RV end systolic and end diastolic volumes, and decreased right ventricular ejection fraction (RVEF) and RV glucose uptake. However, RV mass, SV, and RV free fatty acid uptake remained unchanged in this cohort [171]. In another study that included both men and women participants, HIIT reversed pathological LV remodeling, reduced LV end-diastolic and end-systolic volumes, and improved left ventricular ejection fraction (LVEF) in stable post-infarction heart failure patients; sex differences were not addressed [55]. Furthermore, Wisløff and coworkers postulated that the beneficial impact of HIIT on post-infarction cardiac remodeling may have involved reduction in plasma pro-natriuretic peptide (B-type natriuretic peptide (BNP)) level, a marker of hypertrophy and severity of heart failure [55]. Beneficial cardiovascular effects of HIIT were also seen in aging men. In a study of sedentary aging men and aging male athletes that underwent aerobic preconditioning exercise or regular exercise regimen, respectively, this particular regimen of HIIT training resulted in improved resting blood pressure in both groups without the development of pathological cardiac remodeling [172]. The underlying mechanism(s) of improved cardiac structure and function with HIIT is not clear [156,162,173]. The beneficial effects of HIIT regimen may be attributed to the intermittent periods of rest

that allows recovery, thereby enabling patients to complete the activity while also building their aerobic and anaerobic capacity [55].

Although HIIT is shown to have beneficial effects on cardiac structure and function, reduced ventricular function has also been reported with HIT. In healthy males, high-intensity endurance training caused a greater reduction in global RV strain, which was more pronounced in the RV free wall. Interestingly, a decrease in global LV strain was also observed, which was more pronounced in the LV septum [174]. This observation indicates that HIT may induce segment-specific cardiac dysfunction as well. Detrimental effects of HIT are also seen in the form of cardiac fatigue [8,175,176]. Data are not clear on sex differences in cardiac fatigue with HIT. Cote et al. noted that cardiac fatigue was more pronounced in males compared to females during a HIT (triathlon lasting six hours) [175]. However, sex differences were not noted in cardiac fatigue with prolonged HIT consisting of ultramarathon [175]. Further studies are warranted addressing the sex-specific detrimental effects of HIT.

7.2. Sex Disparity in Vascular Wall Structure and Function with HIT

The beneficial effects of HIIT are also reported on the vasculature. A study by Batacan Jr et al. reported that in adult male rats overfed with a high-fat high carbohydrate (HFHC) diet, HIIT decreased the contractile responses of mesenteric arteries to α -adrenergic stimuli and improved endothelium-dependent vasorelaxation to ACh [177]. These data were promising, as the investigators of this study demonstrated that HIIT was capable of alleviating mesenteric arterial contraction in these male rats despite being on HFHC diet. Females were not included in this study [177]. HIIT also improved vascular wall structure and function in human subjects. HIIT decreased systolic blood pressure in postmenopausal women; males were not studied [178]. In sedentary healthy adults (the sex of the subjects was not indicated), HIIT improved vascular function by increasing FMD of brachial arteries and decreasing aortic PWV, as indicators of endothelial function and arterial wall stiffness, respectively. These results suggest that HIIT alters vascular hemodynamic, thereby decreasing arterial wall thickness and enhancing endothelial function [129]. Similarly, improved FMD of brachial arteries and decreased oxidative stress were reported in the heart failure patients who underwent HIIT. The researchers postulated that the augmented NO bioavailability and plasma antioxidant level might have contributed to the beneficial impacts of HIIT on the vasculature of cardiac failure patients. Both males and females were included in this study, but the investigators did not analyze the sex differences in vascular remodeling [55]. In addition, Rognmo et al. [179] and Freyssin et al. [180] reported that HIIT intervention in both male and female patients with CAD and heart failure was safe and was not associated with adverse cardiac outcomes.

Overall, the data are sparse on the beneficial impacts of HIT on the vasculature, which has been mostly shown to have either detrimental or no effect on the vasculature. For instance, a clinical study conducted only in healthy males showed that HIT did not change the resting brachial artery diameter (a measure of peripheral vessel function) [149]. Furthermore, exclusively in male SHR, HIT adversely altered mesenteric arterial endothelial ultrastructure and function. In the same study, HIT increased oxidative stress and reduced NO bioavailability [181]. Similarly, Chen et al. reported that HIT worsened hypertension and intensified adverse remodeling of L-type voltage-gated Ca^{2+} ($\text{Ca}_v1.2$) channels (up-regulation of $\text{Ca}_v1.2$ channels is a hallmark feature of hypertension) in mesenteric arteries of male SHR; females were not studied [182]. Investigators postulate that the detrimental effects of HIT may be reversed by incorporating HIIT since it exerts potentially greater effects compared to traditional endurance exercise [158]. Overall, HIIT has a more beneficial impact on the cardiovascular system compared to HIT.

Table 4. The described studies associated with the high-intensity training (HIT) regimens.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanisms
Oláh et al. [164]	Healthy male and female Wistar rats	Swimming Duration: 200 min/day, 5 days/week for 12 weeks	↑SV and ↑contractility and stroke work in both sexes	More pronounced LV hypertrophy in females than males, ↑diastolic function only in males	↑Phosphorylation of AKT in the myocardium in both sexes but to a greater degree in females, thus leading to more pronounced LV hypertrophy in females
Verboven et al. [140]	Healthy male Sprague-Dawley rats	Running: Treadmill Duration: 10 bouts, 5 days/week for 13 weeks Speed: 18 m/min at 30° inclination	Beneficial LV hypertrophy, ↑EF, ↑cardiac output and volume, ↓myocardial collagen content, ↑cardiac capillary density	N/A	↑Cardiac metabolism due to increased oxygen supplied by enhanced capillary density and ↑citrate synthase and complex II enzyme activity (measure of mitochondrial mass)
de Oliveira Sá et al. [156]	Male C57BL/6 mice, overfed a diet high in fat or fructose	Running: Treadmill Duration: 3 days/week for 12 weeks Speed: 45 m/min	↓LV mass and LV wall thickness	N/A	Modulated components of the cardiac RAS, ACE2/Angiotensin (1–7)/Mas receptor axis
Brown et al. [161]	Male Sprague Dawley rats with pulmonary arterial hypertension	Running: Treadmill Duration: 30 min/day, 5 times/week for 6 weeks	↓RV systolic pressure, ↓RV hypertrophy, ↓fibrosis, ↑cardiac output	N/A	↑RV apelin expression
Rahimi et al. [170]	Male Wistar rats with IRI	Running: Treadmill Duration: 76–85 min/day, 5 consecutive days	↓Infarct size by 50% and 35% after 1 and 7 days post exercise	N/A	N/A
Batacan Jr et al. [177]	Wistar adult male rats overfed with a high-fat high carbohydrate (HFHC) diet	Running: Treadmill Duration: 4 bouts, 5 days/week for 12 weeks Speed: 50 m/min at 10% inclination	No significant difference for SBP or HR before and after exercise, ↑endothelium-dependent relaxation to acetylcholine, ↓contractile responses of mesenteric arteries to α-adrenergic stimuli	N/A	N/A
Fang et al. [181]	Male SHR	Running: Treadmill Duration: 60 min/day, 5 days/week for 8 weeks Speed: 26–28 m/min (~75–85% of the maximal aerobic velocity)	↑SBP and ↑DBP	N/A	↑Oxidative stress, ↓NO bioavailability
Chen et al. [182]	Male SHR	Running: Treadmill Duration: 60 min/day, 5 days/week for 8 weeks Speed: 26–28 m/min (~75–85% of the maximal aerobic velocity)	Worsened hypertension	N/A	↑Adverse remodeling of L-type voltage-gated Ca ²⁺ (Cav1.2) channels

Table 4. Cont.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanisms
Heiskanen et al. [171]	Healthy, middle-aged males	Cycle ergometer Duration: 6 sessions in 2 weeks HIIT session: 4–6 × 30 s all-out cycling/4 min recovery	Beneficial RV hypertrophy, ↑RV end systolic and end diastolic volumes, ↓RVEF, ↓RV glucose uptake, but RV mass, SV, and RV free fatty acid uptake remained unchanged	N/A	N/A
Stewart et al. [174]	Recreationally active, healthy males who were training >5 h/week	Cycle ergometer Duration: only one 90 min exercise session at 110% of gas exchange threshold (GET)	More pronounced decrements in RV function, ↓LV function only to the sites of septal myocardium	N/A	N/A
Wisløff et al. [55]	Male and female heart failure patients	Walking: Treadmill Duration: 38 min/day, 3 times/week for 12 weeks	Reversed pathological LV remodeling, ↓LV end-diastolic and end-systolic volumes, ↑LVEF, ↑brachial artery FMD	Sex differences were not studied	↓Plasma pro-BNP level, ↑NO bioavailability, ↑Plasma antioxidant level
Grace et al. [172]	Aging male non-athletes and aging male athletes	Sprints, cycle ergometers Duration: Once/5 days for 6 weeks	↑Resting BP in both groups without causing pathological remodeling, ↑diastolic septal thickness, and ↓chamber diameter only in athletes	N/A	N/A
Klonizakis et al. [178]	Postmenopausal females	Cycling Duration: 10 × 1-min intervals at 100% of peak power output, 6 sessions in 2 weeks	↓SBP, no improvement in brachial artery FMD	N/A	N/A
Ramírez-Vélez et al. [129]	Healthy adults (study did not mention the sex of the subjects)	Fast walking and running: Treadmill Duration: 4 × 4 min intervals at 85–95% of HRR, 3 days/week for 12 weeks	↑Brachial artery FMD, ↓aortic PWV	N/A	N/A
Shenouda et al. [149]	Healthy males	Cycling sprints Duration: 3 × 20 s sprint interval training for 10 min for 12 weeks	No change in brachial artery diameter	N/A	N/A

↓decrease, ↑increase, stroke volume (SV), left ventricular (LV), ejection fraction (EF), renin–angiotensin system (RAS), right ventricular (RV), ischemia-reperfusion injury (IRI), systolic blood pressure (SBP), heart rate (HR), spontaneously hypertensive rats (SHR), right ventricular ejection fraction (RVEF), diastolic blood pressure (DBP), nitric oxide (NO), left ventricular ejection fraction (LVEF), flow-mediated dilation (FMD), pro brain natriuretic peptide (pro-BNP), blood pressure (BP), heart rate reserve (HRR), pulse wave velocity (PWV).

7.3. Limitations

The sex-specific impacts of HIT and HIIT on cardiovascular remodeling is not well documented in either animal or human studies. Traditionally, the majority of studies on the impacts of HIT or HIIT have excluded females. Furthermore, those studies that did assess the impacts of HIT on cardiovascular remodeling using both male and female subjects did not attempt to address sex differences in their findings [55,162,179,180], thus preventing the assessment of sex-specific impacts of HIT or HIIT on cardiovascular remodeling. Therefore, the relatively small number of studies that addressed sex differences in the impacts of HIT or HIIT on cardiovascular remodeling limits us from drawing any significant conclusion. Though some studies have shown benefits of HIT on cardiovascular remodeling in CVD, there is a lack of HIT or HIIT studies in healthy subjects. Furthermore, variables, such as strain of animals, human race, age, fluctuations in sex hormones in menstruating females, menopause, diet, and varying HIIT programs (duration, intensity, and frequency) in different studies hamper our ability to assess the impacts of HIT or HIIT on the underlying mechanisms leading to cardiovascular remodeling. Moreover, differences between animal models and humans make it difficult to translate the findings in small-animal models to human studies.

8. Sex-Specific Impact of Combined Exercise on Cardiovascular Remodeling

Combined exercise trains more muscle groups at once in a relatively short period of time. Combined exercise has been demonstrated to offer more cardiovascular benefits in patients with CVD compared to any single exercise modality in several randomized control trials [183–185]. There are different types of combined exercise training, such as aerobic-resistance, aerobic-strength, and endurance-resistance, amongst others. The combination of aerobic and resistance training is frequently used as a combined form of exercise in humans. According to the American College of Sports Medicine (ACSM), aerobic exercise is any activity that increases the capacity of the cardiorespiratory system by increasing oxygen supply and improving the oxygen utilization in the muscles [186]. Examples of aerobic exercise include walking, swimming, dancing, cycling, jogging, and hiking [187]. Resistance training is a type of exercise that includes the use of load, machinery, or your own body weight while exercising the muscles [188]. Strength training and endurance training, however, are subtypes of resistance training. Strength training usually involves a load of 85% or more of one repetition maximum (1RM) for six or less repetitions of two to six sets with rest periods of two to five minutes. Muscular endurance training, on the other hand, requires resistance training at a load of 67% or less of 1RM for twelve or more repetitions of two to three sets with rest periods of thirty seconds or less [189]. We will discuss both animal and human studies addressing the sex disparities in cardiac structure and function with combined exercise below (please refer to Table 5 for details of the exercise regimens).

8.1. Sex Disparity in Cardiac Structure and Function with Combined Exercise

Combined exercise has been shown to improve adverse cardiac remodeling associated with hypertension and aging [190]. In a study conducted only in menopausal female SHR, aerobic-resistance training has been shown to improve cardiac function and decrease heart rates of experimental subjects [191]. Furthermore, Shimojo and coworkers showed that these improvements in the cardiac functions resulted from exercise-augmented cardiovascular autonomic modulation, decreased levels of tumor necrosis factor (TNF) and interleukin-6 (IL-6), reduced nicotinamide adenine dinucleotide phosphate (NADPH) oxidase (one of the major sources of superoxide in the cardiovascular system), and elevated levels of enzymatic or non-enzymatic antioxidants [191].

Clinical studies also concur with the beneficial effects of combined training on cardiac health. A randomized phase III clinical trial that included both male and female patients with heart failure showed that aerobic-strength training improved the LV diastolic function. However, this study did not analyze potential sex-specific impacts of combined exercise [192]. Similarly, in a randomized control trial of post-myocardial infarction (MI) male patients, combined aerobic resistance training showed improvements in LVEF and

diastolic function. Females, however, were not included in this clinical trial [184]. Furthermore, Beckers and coworkers showed that combined endurance-resistance improved the LVEF in male and female patients with CHF, but sex differences were not addressed [183]. In contrast, a meta-analysis conducted in clinically stable male and female CHF patients noted that though aerobic training reversed LV remodeling, this favorable effect was not confirmed when aerobic training was combined with strength training [193]. In the above study, Haykowsky et al. reasoned that lack of improvement in cardiac function when aerobic exercise was combined with strength training could be due to the unfavorable outcome of strength training in cardiac failure patients [193]. Of note, strength training is shown to adversely alter cardiovascular function by increasing systolic and diastolic pressure loading and increasing LV wall stress [193].

8.2. Sex Disparity in Vascular Wall Structure and Function with Combined Exercise

Several studies have demonstrated the beneficial effects of combined exercise on vascular wall structure and function. One study found that combined aerobic-resistance training enhanced the expression of VEGF, an important angiogenic factor, in the skeletal muscle of the fifty-week-old obese male rats. This study did not include female subjects [194]. A study that was conducted exclusively in menopausal SHR showed that aerobic-resistance training decreased mean arterial blood pressure [191]. In this study, Shimojo et al. reported that exercise improved baroreflex sensitivity to increased blood pressure, leading to decreased sympathetic activation of peripheral vessels and reduced arterial blood pressure in those menopausal SHR [191]. Impaired baroreflex sensitivity is common and a strong predictor of arterial hypertension and cardiac mortality [195,196].

Beneficial effects of combined training on vascular function were also noted in human studies. Combined aerobic-resistance training decreased arterial stiffness and blood pressure in hypertensive postmenopausal women [197]. Son et al. reported that the decreased arterial stiffness in hypertensive postmenopausal women may have involved reduced ET-1 and increased NO (as measured by the level of nitrite/nitrate in blood) [197]. Thus, the potential underlying mechanisms for the combined training-mediated improvements in regulation of blood pressure in those subjects were attributed to the enhancement of vascular endothelial function or increased endothelium-dependent vasodilation [197,198]. However, the male subjects were not included in Son et al.'s study. In another study performed in premenopausal sedentary hypertensive women, aerobic-resistance training resulted in significant reduction in blood pressure and heart rate. The investigators of this study suggested that the reduction of vasomotor tone or increased vagal tone may have contributed to beneficial effects on blood pressure and heart rate that occurred after this exercise program [199]. Along the similar lines, Figueroa and coworkers reported that endurance-resistance training decreases blood pressure and arterial stiffness in postmenopausal women; men were not included in this study. These investigators attributed the beneficial effects of combined training on blood pressure to the enhanced endothelium-dependent vasodilation [200].

Studies in the literature showed that high-intensity resistance training might decrease arterial compliance in healthy men. Specifically, resistance training resulted in lowered carotid artery compliance in healthy young men [201]. In healthy middle-aged men (average years of resistance training was 21.3 ± 2.8 years), the carotid artery compliance was also decreased by 30% compared to their sedentary peers [202]. These studies did not include women. However, Kawano et al. demonstrated that resistance training in conjunction with aerobic training improves arterial compliance by decreasing arterial stiffness in healthy men (women were not included) [203]. These findings suggest that combined exercise may ameliorate vascular functions of men and women both in healthy and disease states. However, some controversies exist regarding the beneficial effects of combined exercise. In a previous meta-analysis, combined aerobic-resistance training and isolated aerobic training were reported to have nearly similar effects on arterial stiffness [204]. In contrast, a study performed in hypertensive older adults including both men and women demonstrated that aerobic-resistance training did not show any additional benefits in reducing blood pressure compared to resistance or aerobic training alone [205].

Table 5. The described studies associated with the combined exercise regimens.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanism
Shimojo et al. [191]	Menopausal female SHR	Aerobic: Running on treadmill, 1 h/day at ~50–60% of maximal running speed Resistance: Ladder climbing, 15 climbs/session at 1st–2nd week: 30–40%; 3rd–5th week: 40–50%; and 6th–8th week: 40–60% of the maximal load Duration: 5 days/week for 8 weeks	↓HR, ↓mean arterial BP, ↑baroreflex sensitivity	N/A	↓TNF and IL-6, ↓NADPH oxidase, ↑level of enzymatic or non-enzymatic antioxidants
Chrysohoou et al. [192]	Male and female heart failure patients	Aerobic: Cycle ergometers for 45 min/day Resistance: 4 exercises (knee extension, seated chest press, peck deck and lateral pull-down) with a fitness equipment Duration: 3 days/week for 12 weeks	↓PWV, ↑SBP, ↑LV diastolic function	Sex differences were not studied	N/A
Dor-Haim et al. [184]	Male MI patients	Aerobic-resistance: 20 min of treadmill walking, 15 min of cycling and 10 min of hand cycle paddling, total 45 min/day Duration: Twice/week for 12 weeks	↑LVEF, ↑diastolic function	N/A	N/A
Beckers et al. [183]	Male and female CHF patients	1st–2nd month: 10 min endurance, 40 min resistance 3rd–4th month: 16 min endurance, 30 min resistance 5th–6th month: 10, 12 and 15 min of endurance, the remainder of exercise session spent on resistance training Duration: 70 sessions in 6 months	↑LVEF	Sex differences were not studied	NT-proBNP levels remained unchanged
Son et al. [197]	Postmenopausal hypertensive females	Aerobic-resistance: Exercise intensity was increased gradually from 40% to 70% of HRR/4 weeks Duration: 3 times/week for 12 weeks	↓Brachial-ankle PWV, ↓BP	N/A	↓Endothelin-1, ↑NO (as measured by the level of nitrite/nitrate in blood)
Masroor et al. [199]	Premenopausal hypertensive females	Aerobic: Running on treadmill for 20 min/day at 50–80% of HR _{max} Resistance: 3 sets of 10 repetitions of 5 exercises at an intensity of 50–80% of 1RM Duration: 5 days/week for 4 weeks	↓BP, ↓HR	N/A	N/A

Table 5. Cont.

Study	Participant Characteristics	Exercise Regimen	Cardiovascular Structural and Functional Findings	Sex-Specific Impact	Proposed Molecular Mechanism
Figuerola et al. [200]	Postmenopausal females	Endurance: Walking on treadmill for 20 min at 60% of HR _{max} Resistance: 12 repetitions for 9 exercises for 20 min at 60% of HR _{max} Duration: 3 times/week for 12 weeks	↓Brachial-ankle PWV, ↓SBP and DBP, ↓HR	N/A	N/A
Kawano et al. [203]	Healthy males	Aerobic: Cycling for 30 min at 60% of HR _{max} Resistance: 3 sets of 8–12 exercises at 80% of 1RM Duration: 3 sessions/week for 4 months	↑Arterial compliance	N/A	N/A
Lima et al. [205]	Hypertensive older males and females	Aerobic: Treadmill ergometer, 1st–4th week for 25 min, 5th–10th week for 35 min Resistance: 1st–4th week: 9 exercises, 5th–10th week: 15 repetitions for the upper limbs and 20 repetitions for the trunk and lower limbs at 50 to 60% 1RM Duration: 3 training sessions/week for 10 weeks	Did not show any additional benefits in reducing BP compared to resistance or aerobic training alone	Sex differences were not studied	N/A
Shiotsu et al. [206]	Older males	Aerobic: Cycling for 20 min at 60% of HRR Resistance: 3 sets of 8–12 repetitions for 5 different exercises at 70–80% of 1RM Duration: Twice/week for 10 weeks	↓Carotid-femoral PWV	N/A	N/A
Okamoto et al. [207]	Healthy males and females	Aerobic: Running for 20 min at 60% of the targeted HR Resistance: 5 sets of 8–10 repetitions at 80% of 1RM, Duration: Twice/week for 8 weeks	↓Brachial-ankle PWV, ↑brachial artery FMD	Sex differences were not studied	N/A

↓decrease, ↑increase, spontaneously hypertensive rats (SHR), heart rate (HR), blood pressure (BP), tumor necrosis factor (TNF), interleukin-6 (IL-6), nicotinamide adenine dinucleotide phosphate (NADPH), pulse wave velocity (PWV), systolic blood pressure (SBP), left ventricular (LV), myocardial infarction (MI), left ventricular ejection fraction (LVEF), chronic heart failure (CHF), N-terminal pro b-type natriuretic peptide (NT-proBNP), nitric oxide (NO), maximum heart rate (HR_{max}), diastolic blood pressure (DBP), heart rate reserve (HRR), 1 repetition maximum (1RM), flow-mediated dilation (FMD).

The order of exercise may play an important role on the outcomes of combined exercise. For instance, a study conducted only in older men [206] and another study done in healthy men and women [207] both showed that arterial stiffness was improved when aerobic exercise was performed after high-intensity resistance training, but no benefit was observed when the opposite order was followed. Another study on male and female normotensive and hypertensive adults also showed that aerobic exercise after resistance training exerted either no effects or some beneficial effects on arterial compliance [208]. Shiotsu et al. proposed that aerobic exercise after resistance training promoted arterial flexibility in older men by increasing the production of NO [206]. Moreover, Kawano et al. suggested that when aerobic exercise follows resistance training, it may prevent arterial stiffness associated with resistance training, but this benefit is lost when the exercise order is reversed [203]. None of above studies assessed the potential role of biological sex in relation to the order of exercise on the vascular structure or function.

8.3. Limitations

Sex differences in combined exercise on the cardiovascular remodeling have not been fully explored, as the majority of studies were performed either in males or females. Although some studies included both sexes, they did not isolate or compare the specific role of sex on the outcome of combined exercise. Furthermore, majority of studies were conducted in humans, with a relatively small sample size. Animal studies assessing the benefits of combined exercise on cardiovascular remodeling and functional adaptation are lacking. Moreover, the current literature has focused mainly on the impact of combined exercise on disease state rather than healthy state. With existence of various combinations of aerobic-strength training regimens, it is important to discern which combination is the most beneficial for cardiovascular health based on sex, age, and underlying conditions. Finally, although there are a number of studies that suggest beneficial roles of combined exercise, it should be noted that some studies reported no significant effects on the cardiovascular system [204,205].

9. Conclusion and Future Direction

Given that sex differences play an important role in cardiovascular physiology, the aim of this review was to highlight the studies in the literature that investigated sex-specific benefits of exercise on cardiovascular remodeling. The knowledge gained from current and future studies will ultimately (1) enhance our understanding of the mechanisms underlying the beneficial effects of exercise on cardiovascular function in healthy and disease states, (2) identify the role that biological sex may play in the impact of exercise on restoring cardiovascular function, and (3) provide insight into potential pathways for promoting healthy cardiovascular function and novel therapeutic targets for treating CVD.

Exercise training has emerged as a valuable modality in the prevention of CVD. Since cardiovascular physiology and CVD onset and progression vary between males and females, it is important to assess sex-specific roles of exercise training on the cardiovascular system. Throughout our review, we have underlined several (but not all) studies on the impacts of different types of exercise intensity and their relation to sex on the cardiovascular system. However, the data on sex differences in EICR are limited. Traditionally, studies on the effects of exercise have excluded females. Nevertheless, there has been an increasing number of investigations that have sought to include both sexes. On the other hand, regardless of several studies on EICR in males, there is still lack of consensus in the cardiovascular impacts of androgen in general. Androgens may simultaneously benefit and detriment the cardiovascular system by different mechanisms. Furthermore, most of the literature are focused on the outcomes of exercise in diseased models rather than healthy individuals. Therefore, future studies should be more inclusive of male and female subjects in both healthy and disease states.

In the study of sex-specific impacts of EICR and functional adaptation, it is important to note that there are numerous physiological variables, including age, weight, race, and

body composition (including fat and lean muscle mass), that play roles in EICR. Underlying genetic and molecular differences may also influence the structural remodeling pathways involved in cardiac hypertrophy, inflammation, fibrosis, and apoptosis. Moreover, the fluctuations in the levels of female sex hormone during different phases of the menstrual cycle may impact cardiovascular metabolism and function during exercise. Besides, a majority of the studies on the sex-specific impacts of EICR have not compared the effects of sex steroid hormones (e.g., estrogen vs. progesterone or testosterone vs. dihydrotestosterone (DHT)) on the cardiovascular system. In addition, the effects of female sex hormones signaling in cardiovascular physiology in men or the effects of androgen signaling in women are even less understood. Future investigations should examine the impacts of exercise on the cardiovascular system of the transgender population.

Another possible factor that may impact cardiovascular remodeling is the form of exercise training undertaken; participation in varying types of exercise programs (modality, duration, intensity, and frequency). Future studies should utilize standardized, well-established exercise protocols to assess cardiovascular outcomes in both animal models and human subjects. They also need to assess the long-term effects of EICR and functional adaptation, as most of the studies only focus on short-term outcomes.

In conclusion, exercise has beneficial effects on the cardiovascular system, and it is an important component of health related to the immune system. To date, not many studies have directly addressed the sex-specific roles of different types of exercise on cardiovascular systems in both healthy and disease states. Therefore, it is essential to answer the following questions: (1) Are the cardiovascular protective effects of exercise sex specific? If so, what mechanisms are responsible for the sex-specific impacts of exercise on cardiovascular remodeling? And (2) does biological sex influence the outcome of EICR in disease states?

Overall, the interest in more personalized approaches to the development of selective therapeutic strategies, including exercise, should further advance the studies of sex differences in cardiovascular physiology and pathophysiology. Figure 1 depicts the impacts of exercise and its sex-specific influences on cardiac and vascular remodeling in humans and animals based on studies presented in this review article.

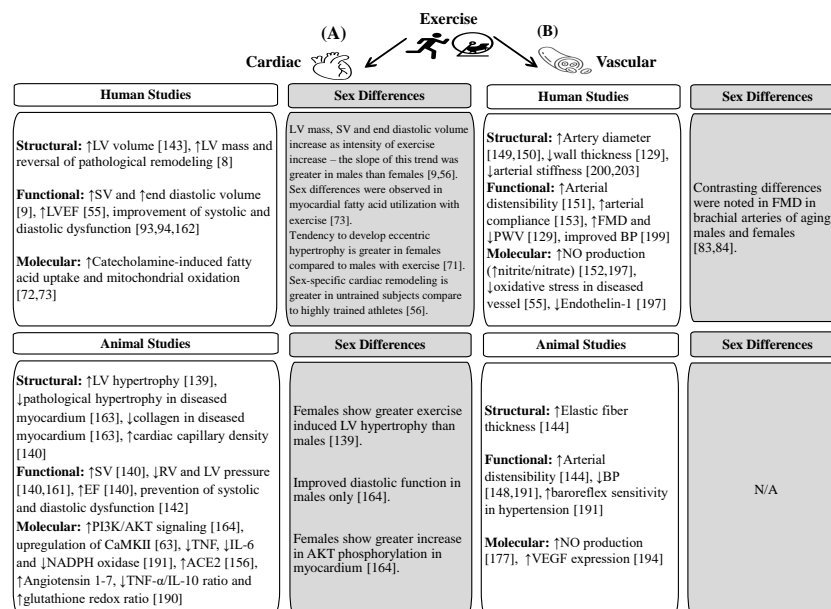


Figure 1. Impacts of exercise and its sex-specific influences on cardiac (A) and vascular (B) remodeling in humans and animals. ↓decrease, ↑increase, left ventricular (LV), stroke volume (SV), left ventricular ejection fraction (LVEF), flow-mediated dilation (FMD), pulse wave velocity (PWV), blood pressure (BP), nitric oxide (NO), right ventricular (RV), ejection fraction (EF), tumor necrosis factor (TNF), interleukin (IL), nicotinamide adenine dinucleotide phosphate (NADPH), angiotensin converting enzyme 2 (ACE2), vascular endothelial growth factor (VEGF).

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References

1. Haram, P.M.; Kemi, O.J.; Wisloff, U. Adaptation of endothelium to exercise training: Insights from experimental studies. *Front. Biosci. J. Virtual Libr.* **2008**, *13*, 336–346. [[CrossRef](#)] [[PubMed](#)]
2. Okada, S.; Hiuge, A.; Makino, H.; Nagumo, A.; Takaki, H.; Konishi, H.; Goto, Y.; Yoshimasa, Y.; Miyamoto, Y. Effect of exercise intervention on endothelial function and incidence of cardiovascular disease in patients with type 2 diabetes. *J. Atheroscler. Thromb.* **2010**, *17*, 828–833. [[CrossRef](#)]
3. Darling, E.A. The Effects of Training. *Boston Med. Surg. J.* **1899**, *141*, 229–233. [[CrossRef](#)]
4. Skidlauf und Skidwettlauf Eine Medizinische Sportstudie. Available online: <https://www.vialibri.net/years/books/98218535/1898-henschen-salomon-eberhard-skidlauf-und-skidwettlauf-skilauf-und-skiwettlauf> (accessed on 10 May 2021).
5. Laughlin, M.H.; Diana, J.N.; Tipton, C.M. Effects of exercise training on coronary reactive hyperemia and blood flow in the dog. *J. Appl. Physiol.* **1978**, *45*, 604–610. [[CrossRef](#)] [[PubMed](#)]
6. Scharhag, J.; Schneider, G.; Urhausen, A.; Rochette, V.; Kramann, B.; Kindermann, W. Athlete’s heart: Right and left ventricular mass and function in male endurance athletes and untrained individuals determined by magnetic resonance imaging. *J. Am. Coll. Cardiol.* **2002**, *40*, 1856–1863. [[CrossRef](#)]
7. Kaptoge, S.; Pennells, L.; Bacquer, D.D.; Cooney, M.T.; Kavousi, M.; Stevens, G.; Riley, L.M.; Savin, S.; Khan, T.; Altay, S.; et al. World Health Organization cardiovascular disease risk charts: Revised models to estimate risk in 21 global regions. *Lancet Glob. Health* **2019**, *7*, e1332–e1345. [[CrossRef](#)]
8. Gielen, S.; Schuler, G.; Adams, V. Cardiovascular effects of exercise training: Molecular mechanisms. *Circulation* **2010**, *122*, 1221–1238. [[CrossRef](#)]
9. Turkbey, E.B.; Jorgensen, N.W.; Johnson, W.C.; Bertoni, A.G.; Polak, J.F.; Diez Roux, A.V.; Tracy, R.P.; Lima, J.A.C.; Bluemke, D.A. Physical activity and physiological cardiac remodelling in a community setting: The Multi-Ethnic Study of Atherosclerosis (MESA). *Heart Br. Card. Soc.* **2010**, *96*, 42–48. [[CrossRef](#)] [[PubMed](#)]
10. Shufelt, C.; Pacheco, C.; Tweet, M.S.; Miller, V.M. Sex-specific physiology and cardiovascular disease. *Adv. Exp. Med. Biol.* **2018**, *1065*, 433–454.
11. dos Santos, R.L.; da Silva, F.B.; Ribeiro, R.F.; Stefanon, I. Sex hormones in the cardiovascular system. *Horm. Mol. Biol. Clin. Investig.* **2014**, *18*, 89–103. [[CrossRef](#)]
12. Barrett-Connor, E. Heart disease in women. *Fertil. Steril.* **1994**, *62*, 127S–132S. [[CrossRef](#)]
13. Rodgers, J.L.; Jones, J.; Bolleddu, S.I.; Vanthenapalli, S.; Rodgers, L.E.; Shah, K.; Karia, K.; Panguluri, S.K. Cardiovascular Risks Associated with Gender and Aging. *J. Cardiovasc. Dev. Dis.* **2019**, *6*, 19. [[CrossRef](#)] [[PubMed](#)]
14. Purkiss, S.; Huckell, V.F. Cardiovascular Physiology: Similarities and Differences between Healthy Women and Men. *J. SOGC* **1997**, *19*, 853–859. [[CrossRef](#)]
15. Wingate, S. Cardiovascular Anatomy and Physiology in the Female. *Crit. Care Nurs. Clin. N. Am.* **1997**, *9*, 447–452. [[CrossRef](#)]
16. de Simone, G.; Devereux, R.B.; Daniels, S.R.; Meyer, R.A. Gender Differences in Left Ventricular Growth. *Hypertension* **1995**, *26*, 979–983. [[CrossRef](#)] [[PubMed](#)]
17. Prabhavathi, K.; Selvi, K.T.; Poornima, K.N.; Sarvanan, A. Role of Biological Sex in Normal Cardiac Function and in its Disease Outcome—A Review. *J. Clin. Diagn. Res. JCDR* **2014**, *8*, BE01–BE04. [[CrossRef](#)] [[PubMed](#)]
18. Cabo, C. The gender inequality of the heart. *Comput. Biol. Med.* **2013**, *43*, 1073–1074. [[CrossRef](#)]
19. Wu, J.; Dai, F.; Li, C.; Zou, Y. Gender Differences in Cardiac Hypertrophy. *J. Cardiovasc. Transl. Res.* **2020**, *13*, 73–84. [[CrossRef](#)]
20. Gilligan, D.M.; Quyyumi, A.A.; Cannon, R.O. Effects of physiological levels of estrogen on coronary vasomotor function in postmenopausal women. *Circulation* **1994**, *89*, 2545–2551. [[CrossRef](#)]
21. Jazbutyte, V.; Arias-Loza, P.A.; Hu, K.; Widder, J.; Govindaraj, V.; von Poser-Klein, C.; Bauersachs, J.; Fritzscheier, K.-H.; Hegele-Hartung, C.; Neyses, L.; et al. Ligand-dependent activation of ER lowers blood pressure and attenuates cardiac hypertrophy in ovariectomized spontaneously hypertensive rats. *Cardiovasc. Res.* **2007**, *77*, 774–781. [[CrossRef](#)]
22. Luo, T.; Kim, J.K. The Role of Estrogen and Estrogen Receptors on Cardiomyocytes: An Overview. *Can. J. Cardiol.* **2016**, *32*, 1017–1025. [[CrossRef](#)] [[PubMed](#)]
23. Pelzer, T.; Jazbutyte, V.; Hu, K.; Segerer, S.; Nahrendorf, M.; Nordbeck, P.; Bonz, A.; Muck, J.; Fritzscheier, K.; Hegelehartung, C. The estrogen receptor- α agonist 16 α -LE2 inhibits cardiac hypertrophy and improves hemodynamic function in estrogen-deficient spontaneously hypertensive rats. *Cardiovasc. Res.* **2005**, *67*, 604–612. [[CrossRef](#)] [[PubMed](#)]
24. Pelzer, T.; Schumann, M.; Neumann, M.; deJager, T.; Stimpel, M.; Serfling, E.; Neyses, L. 17 β -Estradiol Prevents Programmed Cell Death in Cardiac Myocytes. *Biochem. Biophys. Res. Commun.* **2000**, *268*, 192–200. [[CrossRef](#)] [[PubMed](#)]

25. Cavasin, M.A.; Sankey, S.S.; Yu, A.-L.; Menon, S.; Yang, X.-P. Estrogen and testosterone have opposing effects on chronic cardiac remodeling and function in mice with myocardial infarction. *Am. J. Physiol.-Heart Circ. Physiol.* **2003**, *284*, H1560–H1569. [[CrossRef](#)]
26. Papamitsou, T.; Barlagiannis, D.; Papaliagkas, V.; Kotanidou, E.; Dermentzopoulou-Theodoridou, M. Testosterone-induced hypertrophy, fibrosis and apoptosis of cardiac cells—An ultrastructural and immunohistochemical study. *Med. Sci. Monit. Int. Med. J. Exp. Clin. Res.* **2011**, *17*, BR266–BR273. [[CrossRef](#)]
27. Zwadlo, C.; Schmidtmann, E.; Szaroszyk, M.; Kattih, B.; Froese, N.; Hinz, H.; Schmitto, J.D.; Widder, J.; Batkai, S.; Bähre, H.; et al. Antiandrogenic Therapy With Finasteride Attenuates Cardiac Hypertrophy and Left Ventricular Dysfunction. *Circulation* **2015**, *131*, 1071–1081. [[CrossRef](#)]
28. Grandi, A.M.; Venco, A.; Barzizza, F.; Scalise, F.; Pantaleo, P.; Finardi, G. Influence of age and sex on left ventricular anatomy and function in normals. *Cardiology* **1992**, *81*, 8–13. [[CrossRef](#)]
29. Goodale, T.; Sadhu, A.; Petak, S.; Robbins, R. Testosterone and the Heart. *Methodist DeBakey Cardiovasc. J.* **2017**, *13*, 68–72. [[CrossRef](#)]
30. Peterson, L.R.; Soto, P.F.; Herrero, P.; Schechtman, K.B.; Dence, C.; Gropler, R.J. Sex Differences in Myocardial Oxygen and Glucose Metabolism. *J. Nucl. Cardiol. Off. Publ. Am. Soc. Nucl. Cardiol.* **2007**, *14*, 573–581. [[CrossRef](#)]
31. Wittnich, C.; Tan, L.; Wallen, J.; Belanger, M. Sex differences in myocardial metabolism and cardiac function: An emerging concept. *Pflüg. Arch. Eur. J. Physiol.* **2013**, *465*, 719–729. [[CrossRef](#)] [[PubMed](#)]
32. Ahimastos, A.A.; Formosa, M.; Dart, A.M.; Kingwell, B.A. Gender differences in large artery stiffness pre- and post puberty. *J. Clin. Endocrinol. Metab.* **2003**, *88*, 5375–5380. [[CrossRef](#)]
33. Waddell, T.K.; Dart, A.M.; Gatzka, C.D.; Cameron, J.D.; Kingwell, B.A. Women exhibit a greater age-related increase in proximal aortic stiffness than men. *J. Hypertens.* **2001**, *19*, 2205–2212. [[CrossRef](#)] [[PubMed](#)]
34. Martins, D.; Nelson, K.; Pan, D.; Tareen, N.; Norris, K. The effect of gender on age-related blood pressure changes and the prevalence of isolated systolic hypertension among older adults: Data from NHANES III. *J. Gen.-Specif. Med. JGSM Off. J. Partnersh. Womens Health Columbia* **2001**, *4*, 10–13, 20.
35. Berry, K.L.; Cameron, J.D.; Dart, A.M.; Dewar, E.M.; Gatzka, C.D.; Jennings, G.L.; Liang, Y.-L.; Reid, C.M.; Kingwell, B.A. Large-artery stiffness contributes to the greater prevalence of systolic hypertension in elderly women. *J. Am. Geriatr. Soc.* **2004**, *52*, 368–373. [[CrossRef](#)] [[PubMed](#)]
36. Rebecca, S.; Matthew, C.Y.; Adam, D.G.; Claudia, E.K.; Joel, K.; Laura, A.C.; Kiang, L.; James, S.H. Sex Differences in Predictors of Longitudinal Changes in Carotid Artery Stiffness. *Arterioscler. Thromb. Vasc. Biol.* **2015**, *35*, 478–484.
37. Mendelsohn, M.E.; Karas, R.H. The protective effects of estrogen on the cardiovascular system. *N. Engl. J. Med.* **1999**, *340*, 1801–1811. [[CrossRef](#)]
38. Rahimian, R.; Dubé, G.P.; Toma, W.; Dos Santos, N.; McManus, B.M.; van Breemen, C. Raloxifene enhances nitric oxide release in rat aorta via increasing endothelial nitric oxide mRNA expression. *Eur. J. Pharmacol.* **2002**, *434*, 141–149. [[CrossRef](#)]
39. Rahimian, R.; Laher, I.; Dube, G.; van Breemen, C. Estrogen and selective estrogen receptor modulator LY117018 enhance release of nitric oxide in rat aorta. *J. Pharmacol. Exp. Ther.* **1997**, *283*, 116–122.
40. Rahimian, R.; Van Breemen, C.; Karkan, D.; Dube, G.; Laher, I. Estrogen augments cyclopiazonic acid-mediated, endothelium-dependent vasodilation. *Eur. J. Pharmacol.* **1997**, *327*, 143–149. [[CrossRef](#)]
41. Stampfer, M.J.; Colditz, G.A.; Willett, W.C.; Manson, J.E.; Rosner, B.; Speizer, F.E.; Hennekens, C.H. Postmenopausal estrogen therapy and cardiovascular disease. Ten-year follow-up from the nurses' health study. *N. Engl. J. Med.* **1991**, *325*, 756–762. [[CrossRef](#)]
42. Reis, S.E.; Gloth, S.T.; Blumenthal, R.S.; Resar, J.R.; Zacur, H.A.; Gerstenblith, G.; Brinker, J.A. Ethinyl estradiol acutely attenuates abnormal coronary vasomotor responses to acetylcholine in postmenopausal women. *Circulation* **1994**, *89*, 52–60. [[CrossRef](#)]
43. Hayashi, T.; Fukuto, J.M.; Ignarro, L.J.; Chaudhuri, G. Basal release of nitric oxide from aortic rings is greater in female rabbits than in male rabbits: Implications for atherosclerosis. *Proc. Natl. Acad. Sci. USA* **1992**, *89*, 11259–11263. [[CrossRef](#)]
44. Kähönen, M.; Tolvanen, J.P.; Sallinen, K.; Wu, X.; Pörsti, I. Influence of gender on control of arterial tone in experimental hypertension. *Am. J. Physiol.* **1998**, *275*, H15–H22. [[CrossRef](#)] [[PubMed](#)]
45. Crews, J.K.; Khalil, R.A. Gender-specific inhibition of Ca²⁺ entry mechanisms of arterial vasoconstriction by sex hormones. *Clin. Exp. Pharmacol. Physiol.* **1999**, *26*, 707–715. [[CrossRef](#)] [[PubMed](#)]
46. Han, X.; Shaligram, S.; Zhang, R.; Anderson, L.; Rahimian, R. Sex-specific vascular responses of the rat aorta: Effects of moderate term (intermediate stage) streptozotocin-induced diabetes. *Can. J. Physiol. Pharmacol.* **2016**, *94*, 408–415. [[CrossRef](#)]
47. Stallone, J.N.; Crofton, J.T.; Share, L. Sexual dimorphism in vasopressin-induced contraction of rat aorta. *Am. J. Physiol.* **1991**, *260*, H453–H458. [[CrossRef](#)] [[PubMed](#)]
48. Tostes, R.C.; David, F.L.; Carvalho, M.H.; Nigro, D.; Scivoletto, R.; Fortes, Z.B. Gender differences in vascular reactivity to endothelin-1 in deoxycorticosterone-salt hypertensive rats. *J. Cardiovasc. Pharmacol.* **2000**, *36*, S99–S101. [[CrossRef](#)]
49. Foryst-Ludwig, A.; Kintscher, U. Sex differences in exercise-induced cardiac hypertrophy. *Pflugers Arch.* **2013**, *465*, 731–737. [[CrossRef](#)] [[PubMed](#)]
50. Baggish, A.L.; Wang, F.; Weiner, R.B.; Elinoff, J.M.; Tournoux, F.; Boland, A.; Picard, M.H.; Hutter, A.M.; Wood, M.J. Training-specific changes in cardiac structure and function: A prospective and longitudinal assessment of competitive athletes. *J. Appl. Physiol. Bethesda Md.* **2008**, *104*, 1121–1128. [[CrossRef](#)] [[PubMed](#)]

51. Caspersen, C.J.; Powell, K.E.; Christenson, G.M. Physical activity, exercise, and physical fitness: Definitions and distinctions for health-related research. *Public Health Rep.* **1985**, *100*, 126–131.
52. de Mendes, M.A.; da Silva, I.; Ramires, V.; Reichert, F.; Martins, R.; Ferreira, R.; Tomasi, E. Metabolic equivalent of task (METs) thresholds as an indicator of physical activity intensity. *PLoS ONE* **2018**, *13*, e0200701. [[CrossRef](#)]
53. Boutcher, Y.N.; Boutcher, S.H. Exercise intensity and hypertension: What's new? *J. Hum. Hypertens.* **2017**, *31*, 157–164. [[CrossRef](#)]
54. Liao, J.; Li, Y.; Zeng, F.; Wu, Y. Regulation of mTOR Pathway in Exercise-induced Cardiac Hypertrophy. *Int. J. Sports Med.* **2015**, *36*, 343–350. [[CrossRef](#)]
55. Wisløff, U.; Støylen, A.; Loennechen, J.P.; Bruvold, M.; Rognum, Ø.; Haram, P.M.; Tjønnå, A.E.; Helgerud, J.; Slørdahl, S.A.; Lee, S.J.; et al. Superior Cardiovascular Effect of Aerobic Interval Training Versus Moderate Continuous Training in Heart Failure Patients: A Randomized Study. *Circulation* **2007**, *115*, 3086–3094. [[CrossRef](#)]
56. Barnes, J.N.; Fu, Q. Sex-Specific Ventricular and Vascular Adaptations to Exercise. *Adv. Exp. Med. Biol.* **2018**, *1065*, 329–346. [[PubMed](#)]
57. Charkoudian, N.; Joyner, M.J. Physiologic considerations for exercise performance in women. *Clin. Chest Med.* **2004**, *25*, 247–255. [[CrossRef](#)] [[PubMed](#)]
58. Wiebe, C.G.; Gledhill, N.; Warburton, D.E.; Jamnik, V.K.; Ferguson, S. Exercise cardiac function in endurance-trained males versus females. *Clin. J. Sport Med. Off. J. Can. Acad. Sport Med.* **1998**, *8*, 272–279. [[CrossRef](#)] [[PubMed](#)]
59. Muhl, C.; Dassen, W.R.M.; Kuipers, H. Cardiac remodelling: Concentric versus eccentric hypertrophy in strength and endurance athletes. *Neth. Heart J.* **2008**, *16*, 129–133. [[CrossRef](#)]
60. Foryst-Ludwig, A.; Kreissl, M.C.; Sprang, C.; Thalke, B.; Böhm, C.; Benz, V.; Gürgen, D.; Dragun, D.; Schubert, C.; Mai, K.; et al. Sex differences in physiological cardiac hypertrophy are associated with exercise-mediated changes in energy substrate availability. *Am. J. Physiol. Heart Circ. Physiol.* **2011**, *301*, H115–H122. [[CrossRef](#)] [[PubMed](#)]
61. Haines, C.D.; Harvey, P.A.; Leinwand, L.A. Estrogens mediate cardiac hypertrophy in a stimulus-dependent manner. *Endocrinology* **2012**, *153*, 4480–4490. [[CrossRef](#)]
62. Konhilas, J.P.; Maass, A.H.; Luckey, S.W.; Stauffer, B.L.; Olson, E.N.; Leinwand, L.A. Sex modifies exercise and cardiac adaptation in mice. *Am. J. Physiol. Heart Circ. Physiol.* **2004**, *287*, H2768–H2776. [[CrossRef](#)] [[PubMed](#)]
63. Luczak, E.D.; Leinwand, L.A. Sex-Based Cardiac Physiology. *Annu. Rev. Physiol.* **2009**, *71*, 1–18. [[CrossRef](#)]
64. Dahlöf, B. Effect of angiotensin II blockade on cardiac hypertrophy and remodelling: A review. *J. Hum. Hypertens.* **1995**, *9* (Suppl. 5), S37–S44.
65. Santos, R.A.S.; Ferreira, A.J. Angiotensin-(1–7) and the renin-angiotensin system. *Curr. Opin. Nephrol. Hypertens.* **2007**, *16*, 122–128. [[CrossRef](#)] [[PubMed](#)]
66. Guimarães, G.G.; Santos, S.H.S.; Oliveira, M.L.; Pimenta-Velloso, E.P.; Motta, D.F.; Martins, A.S.; Alenina, N.; Bader, M.; Santos, R.A.S.; Campagnole-Santos, M.J. Exercise induces renin-angiotensin system unbalance and high collagen expression in the heart of Mas-deficient mice. *Peptides* **2012**, *38*, 54–61. [[CrossRef](#)]
67. Botelho-Santos, G.A.; Sampaio, W.O.; Reudelhuber, T.L.; Bader, M.; Campagnole-Santos, M.J.; Souza dos Santos, R.A. Expression of an angiotensin-(1-7)-producing fusion protein in rats induced marked changes in regional vascular resistance. *Am. J. Physiol. Heart Circ. Physiol.* **2007**, *292*, H2485–H2490. [[CrossRef](#)] [[PubMed](#)]
68. Castro, C.H.; Santos, R.A.S.; Ferreira, A.J.; Bader, M.; Alenina, N.; Almeida, A.P. Effects of genetic deletion of angiotensin-(1-7) receptor Mas on cardiac function during ischemia/reperfusion in the isolated perfused mouse heart. *Life Sci.* **2006**, *80*, 264–268. [[CrossRef](#)]
69. Sniderman, J.D.S.; Sado, D.M.; Sniderman, A.D.; McKenna, W.J. Evaluation of Suspected Right Ventricular Pathology in the Athlete. *Prog. Cardiovasc. Dis.* **2012**, *54*, 397–406. [[CrossRef](#)]
70. Pelliccia, A. Assessment of Left Ventricular Hypertrophy in a Trained Athlete: Differential Diagnosis of Physiologic Athlete's Heart From Pathologic Hypertrophy. *Prog. Cardiovasc. Dis.* **2012**, *10*, 387–396. [[CrossRef](#)]
71. Colombo, C.S.S.S.; Finocchiaro, G. The Female Athlete's Heart: Facts and Fallacies. *Curr. Treat. Options Cardiovasc. Med.* **2018**, *20*, 1–14. [[CrossRef](#)] [[PubMed](#)]
72. Dorn Gerald, W. The Fuzzy Logic of Physiological Cardiac Hypertrophy. *Hypertension* **2007**, *49*, 962–970. [[CrossRef](#)]
73. Soto, P.F.; Herrero, P.; Schechtman, K.B.; Waggoner, A.D.; Baumstark, J.M.; Ehsani, A.A.; Gropler, R.J. Exercise training impacts the myocardial metabolism of older individuals in a gender-specific manner. *Am. J. Physiol. Heart Circ. Physiol.* **2008**, *295*, H842–H850. [[CrossRef](#)] [[PubMed](#)]
74. Iskandar, A.; Mujtaba, M.T.; Thompson, P. Left Atrium Size in Elite Athletes. *JACC Cardiovasc. Imaging* **2015**, *8*, 753–762. [[CrossRef](#)]
75. Król, W.; Jędrzejewska, I.; Konopka, M.; Burkhard-Jagodźńska, K.; Klusiewicz, A.; Pokrywka, A.; Chwalbińska, J.; Sitkowski, D.; Dłużniewski, M.; Mamcarz, A.; et al. Left Atrial Enlargement in Young High-Level Endurance Athletes—Another Sign of Athlete's Heart? *J. Hum. Kinet.* **2016**, *53*, 81–90. [[CrossRef](#)]
76. D'Andrea, A.; Riegler, L.; Golia, E.; Cocchia, R.; Scarafilo, R.; Salerno, G.; Pezzullo, E.; Nunziata, L.; Citro, R.; Cuomo, S.; et al. Range of right heart measurements in top-level athletes: The training impact. *Int. J. Cardiol.* **2013**, *164*, 48–57. [[CrossRef](#)]
77. Yoon, H.J.; Kim, K.H.; Hornsby, K.; Park, J.H.; Park, H.; Kim, H.Y.; Cho, J.Y.; Ahn, Y.; Jeong, M.H.; Cho, J.G. Gender Difference of Cardiac Remodeling in University Athletes: Results from 2015 Gwangju Summer Universiade. *Korean Circ. J.* **2021**, *51*, e34. [[CrossRef](#)] [[PubMed](#)]

78. Green, D.J.; Maiorana, A.; O'Driscoll, G.; Taylor, R. Effect of exercise training on endothelium-derived nitric oxide function in humans. *J. Physiol.* **2004**, *561*, 1–25. [[CrossRef](#)] [[PubMed](#)]
79. Otsuki, T.; Maeda, S.; Iemitsu, M.; Saito, Y.; Tanimura, Y.; Ajisaka, R.; Miyauchi, T. Vascular endothelium-derived factors and arterial stiffness in strength- and endurance-trained men. *Am. J. Physiol.-Heart Circ. Physiol.* **2007**, *292*, H786–H791. [[CrossRef](#)] [[PubMed](#)]
80. Prior, B.M.; Lloyd, P.G.; Yang, H.T.; Terjung, R.L. Exercise-induced vascular remodeling. *Exerc. Sport Sci. Rev.* **2003**, *31*, 26–33. [[CrossRef](#)]
81. Moncada, S.; Palmer, R.M.; Higgs, E.A. Nitric oxide: Physiology, pathophysiology, and pharmacology. *Pharmacol. Rev.* **1991**, *43*, 109–142.
82. Esposito, F.; Mathieu-Costello, O.; Entin, P.L.; Wagner, P.D.; Richardson, R.S. The skeletal muscle VEGF mRNA response to acute exercise in patients with chronic heart failure. *Growth Factors Chur Switz.* **2010**, *28*, 139–147. [[CrossRef](#)] [[PubMed](#)]
83. Black, M.A.; Cable, N.T.; Thijssen, D.H.J.; Green, D.J. Impact of age, sex, and exercise on brachial artery flow-mediated dilatation. *Am. J. Physiol. Heart Circ. Physiol.* **2009**, *297*, H1109–H1116. [[CrossRef](#)] [[PubMed](#)]
84. Pierce, G.L.; Eskurza, I.; Walker, A.E.; Fay, T.N.; Seals, D.R. Sex-specific effects of habitual aerobic exercise on brachial artery flow-mediated dilation in middle-aged and older adults. *Clin. Sci. Lond. Engl.* **2011**, *120*, 13–23. [[CrossRef](#)]
85. Tinken, T.M.; Thijssen, D.H.J.; Hopkins, N.; Dawson, E.A.; Cable, N.T.; Green, D.J. Shear stress mediates endothelial adaptations to exercise training in humans. *Hypertens. Dallas Tex* **2010**, *55*, 312–318. [[CrossRef](#)]
86. Moreau, K.L.; Stauffer, B.L.; Kohrt, W.M.; Seals, D.R. Essential Role of Estrogen for Improvements in Vascular Endothelial Function With Endurance Exercise in Postmenopausal Women. *J. Clin. Endocrinol. Metab.* **2013**, *98*, 4507–4515. [[CrossRef](#)]
87. Ascensão, A.; Ferreira, R.; Magalhães, J. Exercise-induced cardioprotection—biochemical, morphological and functional evidence in whole tissue and isolated mitochondria. *Int. J. Cardiol.* **2007**, *117*, 16–30. [[CrossRef](#)]
88. Powers, S.K.; Demirel, H.A.; Vincent, H.K.; Coombes, J.S.; Naito, H.; Hamilton, K.L.; Shanely, R.A.; Jessup, J. Exercise training improves myocardial tolerance to in vivo ischemia-reperfusion in the rat. *Am. J. Physiol.* **1998**, *275*, R1468–R1477. [[CrossRef](#)]
89. Powers, S.K.; Quindry, J.C.; Kavazis, A.N. Exercise-induced cardioprotection against myocardial ischemia-reperfusion injury. *Free Radic. Biol. Med.* **2008**, *44*, 193–201. [[CrossRef](#)]
90. Hamilton, K.L.; Quindry, J.C.; French, J.P.; Staib, J.; Hughes, J.; Mehta, J.L.; Powers, S.K. MnSOD antisense treatment and exercise-induced protection against arrhythmias. *Free Radic. Biol. Med.* **2004**, *37*, 1360–1368. [[CrossRef](#)]
91. Kanter, M.M.; Hamlin, R.L.; Unverferth, D.V.; Davis, H.W.; Merola, A.J. Effect of exercise training on antioxidant enzymes and cardiotoxicity of doxorubicin. *J. Appl. Physiol. Bethesda Md.* **1985**, *59*, 1298–1303. [[CrossRef](#)] [[PubMed](#)]
92. Powers, S.K.; Criswell, D.; Lawler, J.; Martin, D.; Lieu, F.K.; Ji, L.L.; Herb, R.A. Rigorous exercise training increases superoxide dismutase activity in ventricular myocardium. *Am. J. Physiol.* **1993**, *265*, H2094–H2098. [[CrossRef](#)]
93. Giannuzzi, P.; Temporelli, P.L.; Corrà, U.; Tavazzi, L. ELVD-CHF Study Group Antiremodeling effect of long-term exercise training in patients with stable chronic heart failure: Results of the Exercise in Left Ventricular Dysfunction and Chronic Heart Failure (ELVD-CHF) Trial. *Circulation* **2003**, *108*, 554–559. [[CrossRef](#)] [[PubMed](#)]
94. Hambrecht, R.; Gielen, S.; Linke, A.; Fiehn, E.; Yu, J.; Walther, C.; Schoene, N.; Schuler, G. Effects of exercise training on left ventricular function and peripheral resistance in patients with chronic heart failure: A randomized trial. *JAMA* **2000**, *283*, 3095–3101. [[CrossRef](#)] [[PubMed](#)]
95. McMullen, J.R.; Amirahmadi, F.; Woodcock, E.A.; Schinke-Braun, M.; Bouwman, R.D.; Hewitt, K.A.; Mollica, J.P.; Zhang, L.; Zhang, Y.; Shioi, T.; et al. Protective effects of exercise and phosphoinositide 3-kinase(p110alpha) signaling in dilated and hypertrophic cardiomyopathy. *Proc. Natl. Acad. Sci. USA* **2007**, *104*, 612–617. [[CrossRef](#)] [[PubMed](#)]
96. Lu, L.; Mei, D.F.; Gu, A.-G.; Wang, S.; Lentzner, B.; Gutstein, D.E.; Zwas, D.; Homma, S.; Yi, G.-H.; Wang, J. Exercise training normalizes altered calcium-handling proteins during development of heart failure. *J. Appl. Physiol. Bethesda Md.* **2002**, *92*, 1524–1530. [[CrossRef](#)] [[PubMed](#)]
97. Rolim, N.P.L.; Medeiros, A.; Rosa, K.T.; Mattos, K.C.; Irigoyen, M.C.; Krieger, E.M.; Krieger, J.E.; Negrão, C.E.; Brum, P.C. Exercise training improves the net balance of cardiac Ca²⁺ handling protein expression in heart failure. *Physiol. Genomics* **2007**, *29*, 246–252. [[CrossRef](#)]
98. Coats, A.J.; Adamopoulos, S.; Radaelli, A.; McCance, A.; Meyer, T.E.; Bernardi, L.; Solda, P.L.; Davey, P.; Ormerod, O.; Forfar, C. Controlled trial of physical training in chronic heart failure. Exercise performance, hemodynamics, ventilation, and autonomic function. *Circulation* **1992**, *85*, 2119–2131. [[CrossRef](#)]
99. Braith, R.W.; Welsch, M.A.; Feigenbaum, M.S.; Kluess, H.A.; Pepine, C.J. Neuroendocrine activation in heart failure is modified by endurance exercise training. *J. Am. Coll. Cardiol.* **1999**, *34*, 1170–1175. [[CrossRef](#)]
100. Xu, X.; Wan, W.; Powers, A.S.; Li, J.; Ji, L.; Lao, S.; Wilson, B.; Erikson, J.M.; Zhang, J.Q. Effects of exercise training on cardiac function and myocardial remodeling in post myocardial infarction rats. *J. Mol. Cell. Cardiol.* **2008**, *44*, 114–122. [[CrossRef](#)]
101. Hambrecht, R.; Adams, V.; Erbs, S.; Linke, A.; Kränkel, N.; Shu, Y.; Baither, Y.; Gielen, S.; Thiele, H.; Gummert, J.F.; et al. Regular physical activity improves endothelial function in patients with coronary artery disease by increasing phosphorylation of endothelial nitric oxide synthase. *Circulation* **2003**, *107*, 3152–3158. [[CrossRef](#)]
102. Adams, V.; Linke, A.; Kränkel, N.; Erbs, S.; Gielen, S.; Möbius-Winkler, S.; Gummert, J.F.; Mohr, F.W.; Schuler, G.; Hambrecht, R. Impact of regular physical activity on the NAD(P)H oxidase and angiotensin receptor system in patients with coronary artery disease. *Circulation* **2005**, *111*, 555–562. [[CrossRef](#)]

103. Elliott, A.D.; La Gerche, A. The right ventricle following prolonged endurance exercise: Are we overlooking the more important side of the heart? A meta-analysis. *Br. J. Sports Med.* **2015**, *49*, 724–729. [[CrossRef](#)]
104. La Gerche, A.; Claessen, G. Is Exercise Good for the Right Ventricle? Concepts for Health and Disease. *Can. J. Cardiol.* **2015**, *31*, 502–508. [[CrossRef](#)]
105. Lewis, G.D.; Bossone, E.; Naeije, R.; Grünig, E.; Saggari, R.; Lancellotti, P.; Ghio, S.; Varga, J.; Rajagopalan, S.; Oudiz, R.; et al. Pulmonary Vascular Hemodynamic Response to Exercise in Cardiopulmonary Diseases. *Circulation* **2013**, *128*, 1470–1479. [[CrossRef](#)]
106. Naeije, R.; Chesler, N. Pulmonary Circulation at Exercise. *Compr. Physiol.* **2012**, *2*, 711–741.
107. La Gerche, A.; Roberts, T.; Claessen, G. The response of the pulmonary circulation and right ventricle to exercise: Exercise-induced right ventricular dysfunction and structural remodeling in endurance athletes (2013 Grover Conference series). *Pulm. Circ.* **2014**, *4*, 407–416. [[CrossRef](#)]
108. Finocchiaro, G.; Papadakis, M.; Robertus, J.-L.; Dhutia, H.; Steriotis, A.K.; Tome, M.; Mellor, G.; Merghani, A.; Malhotra, A.; Behr, E.; et al. Etiology of Sudden Death in Sports: Insights From a United Kingdom Regional Registry. *J. Am. Coll. Cardiol.* **2016**, *67*, 2108–2115. [[CrossRef](#)] [[PubMed](#)]
109. Rich, L.; Rarick, J.; Prahlow, J. Arrhythmogenic Right Ventricular Cardiomyopathy in a Young Athlete. *Am. J. Forensic Med. Pathol.* **2021**, *42*, 64–66. [[CrossRef](#)] [[PubMed](#)]
110. Gasperetti, A.; James, C.A.; Cerrone, M.; Delmar, M.; Calkins, H.; Duru, F. Arrhythmogenic right ventricular cardiomyopathy and sports activity: From molecular pathways in diseased hearts to new insights into the athletic heart mimicry. *Eur. Heart J.* **2021**, *42*, 1231–1243. [[CrossRef](#)] [[PubMed](#)]
111. Guasch, E.; Mont, L. Diagnosis, pathophysiology, and management of exercise-induced arrhythmias. *Nat. Rev. Cardiol.* **2017**, *14*, 88–101. [[CrossRef](#)] [[PubMed](#)]
112. Flannery, M.D.; Kalman, J.M.; Sanders, P.; La Gerche, A. State of the Art Review: Atrial Fibrillation in Athletes. *Heart Lung Circ.* **2017**, *26*, 983–989. [[CrossRef](#)]
113. Wilhelm, M.; Roten, L.; Tanner, H.; Wilhelm, I.; Schmid, J.-P.; Saner, H. Gender Differences of Atrial and Ventricular Remodeling and Autonomic Tone in Nonelite Athletes. *Am. J. Cardiol.* **2011**, *108*, 1489–1495. [[CrossRef](#)] [[PubMed](#)]
114. Chhabra, L.; Devadoss, R.; Chaubey, V.K.; Spodick, D.H. Interatrial Block in the Modern Era. *Curr. Cardiol. Rev.* **2014**, *10*, 181–189. [[CrossRef](#)] [[PubMed](#)]
115. Guasch, E.; Benito, B.; Qi, X.; Cifelli, C.; Naud, P.; Shi, Y.; Mighiu, A.; Tardif, J.-C.; Tadevosyan, A.; Chen, Y.; et al. Atrial fibrillation promotion by endurance exercise: Demonstration and mechanistic exploration in an animal model. *J. Am. Coll. Cardiol.* **2013**, *62*, 68–77. [[CrossRef](#)] [[PubMed](#)]
116. Lehmann, M.H.; Timothy, K.W.; Frankovich, D.; Fromm, B.S.; Keating, M.; Locati, E.H.; Taggart, R.T.; Towbin, J.A.; Moss, A.J.; Schwartz, P.J.; et al. Age-Gender Influence on the Rate-Corrected QT Interval and the QT-Heart Rate Relation in Families With Genotypically Characterized Long QT Syndrome. *J. Am. Coll. Cardiol.* **1997**, *29*, 93–99. [[CrossRef](#)]
117. Locati, E.H.; Zareba, W.; Moss, A.J.; Schwartz, P.J.; Vincent, G.M.; Lehmann, M.H.; Towbin, J.A.; Priori, S.G.; Napolitano, C.; Robinson, J.L.; et al. Age- and Sex-Related Differences in Clinical Manifestations in Patients With Congenital Long-QT Syndrome. *Circulation* **1998**, *97*, 2237–2244. [[CrossRef](#)]
118. Benito, B.; Sarkozy, A.; Mont, L.; Henkens, S.; Berruezo, A.; Tamborero, D.; Arzamendi, D.; Berne, P.; Brugada, R.; Brugada, P.; et al. Gender Differences in Clinical Manifestations of Brugada Syndrome. *J. Am. Coll. Cardiol.* **2008**, *52*, 1567–1573. [[CrossRef](#)]
119. Vlachos, K.; Mascia, G.; Martin, C.A.; Bazoukis, G.; Frontera, A.; Cheniti, G.; Letsas, K.P.; Efremidis, M.; Georgopoulos, S.; Gkalapis, C.; et al. Atrial fibrillation in Brugada syndrome: Current perspectives. *J. Cardiovasc. Electrophysiol.* **2020**, *31*, 975–984. [[CrossRef](#)]
120. Mascia, G.; Bona, R.D.; Ameri, P.; Canepa, M.; Porto, I.; Parati, G.; Crotti, L.; Brignole, M. Brugada syndrome and syncope: A practical approach for diagnosis and treatment. *EP Eur.* **2021**, *23*, 996–1002.
121. Saarel, E.V.; Law, I.; Berul, C.I.; Ackerman, M.J.; Kanter, R.J.; Sanatani, S.; Cohen, M.I.; Berger, S.; Fischbach, P.S.; Burton, D.A.; et al. Safety of Sports for Young Patients With Implantable Cardioverter-Defibrillators. *Circ. Arrhythm. Electrophysiol.* **2018**, *11*, e006305. [[CrossRef](#)]
122. Lampert, R. Sport Participation in Patients with Implantable Cardioverter-Defibrillators. *Curr. Treat. Options Cardiovasc. Med.* **2019**, *21*, 66. [[CrossRef](#)] [[PubMed](#)]
123. Lehnart, S.E.; Wehrens, X.H.T.; Laitinen, P.J.; Reiken, S.R.; Deng, S.-X.; Cheng, Z.; Landry, D.W.; Kontula, K.; Swan, H.; Marks, A.R. Sudden Death in Familial Polymorphic Ventricular Tachycardia Associated With Calcium Release Channel (Ryanodine Receptor) Leak. *Circulation* **2004**, *109*, 3208–3214. [[CrossRef](#)] [[PubMed](#)]
124. Noseworthy, P.A.; Tikkanen, J.T.; Porthan, K.; Oikarinen, L.; Pietila, A.; Harald, K.; Peloso, G.M.; Merchant, F.M.; Jula, A.; Väänänen, H.; et al. The early repolarization pattern in the general population: Clinical correlates and heritability. *J. Am. Coll. Cardiol.* **2011**, *57*, 2284–2289. [[CrossRef](#)]
125. Tester, D.J.; Arya, P.; Will, M.; Haglund, C.M.; Farley, A.L.; Makielski, J.C.; Ackerman, M.J. Genotypic heterogeneity and phenotypic mimicry among unrelated patients referred for catecholaminergic polymorphic ventricular tachycardia genetic testing. *Heart Rhythm* **2006**, *3*, 800–805. [[CrossRef](#)]
126. Adams, V. CrossTalk proposal: Acute exercise elicits damage to the endothelial layer of systemic blood vessels in healthy individuals. *J. Physiol.* **2018**, *596*, 537–539. [[CrossRef](#)]

127. Dawson, E.A.; Whyte, G.P.; Black, M.A.; Jones, H.; Hopkins, N.; Oxborough, D.; Gaze, D.; Shave, R.E.; Wilson, M.; George, K.P.; et al. Changes in vascular and cardiac function after prolonged strenuous exercise in humans. *J. Appl. Physiol. Bethesda Md.* **2008**, *105*, 1562–1568. [[CrossRef](#)] [[PubMed](#)]
128. Birk, G.K.; Dawson, E.A.; Batterham, A.M.; Atkinson, G.; Cable, T.; Thijssen, D.H.J.; Green, D.J. Effects of exercise intensity on flow mediated dilation in healthy humans. *Int. J. Sports Med.* **2013**, *34*, 409–414. [[CrossRef](#)]
129. Ramírez-Vélez, R.; Hernández-Quñones, P.A.; Tordecilla-Sanders, A.; Álvarez, C.; Ramírez-Campillo, R.; Izquierdo, M.; Correa-Bautista, J.E.; Garcia-Hermoso, A.; Garcia, R.G. Effectiveness of HIIT compared to moderate continuous training in improving vascular parameters in inactive adults. *Lipids Health Dis.* **2019**, *18*, 1–10. [[CrossRef](#)]
130. Johnson, B.D.; Padilla, J.; Wallace, J.P. The exercise dose affects oxidative stress and brachial artery flow-mediated dilation in trained men. *Eur. J. Appl. Physiol.* **2012**, *112*, 33–42. [[CrossRef](#)]
131. O'Sullivan, S.E. The effects of exercise training on markers of endothelial function in young healthy men. *Int. J. Sports Med.* **2003**, *24*, 404–409.
132. Uhlemann, M.; Möbius-Winkler, S.; Fikenzer, S.; Adam, J.; Redlich, M.; Möhlenkamp, S.; Hilberg, T.; Schuler, G.C.; Adams, V. Circulating microRNA-126 increases after different forms of endurance exercise in healthy adults. *Eur. J. Prev. Cardiol.* **2014**, *21*, 484–491. [[CrossRef](#)]
133. Boos, C.J.; Balakrishnan, B.; Lip, G.Y.H. The effects of exercise stress testing on soluble E-selectin, von Willebrand factor, and circulating endothelial cells as indices of endothelial damage/dysfunction. *Ann. Med.* **2008**, *40*, 66–73. [[CrossRef](#)]
134. Sapp, R.M.; Hagberg, J.M. CrossTalk opposing view: Acute exercise does not elicit damage to the endothelial layer of systemic blood vessels in healthy individuals. *J. Physiol.* **2018**, *596*, 541–544. [[CrossRef](#)]
135. Ghadieh, A.S.; Saab, B. Evidence for exercise training in the management of hypertension in adults. *Can. Fam. Physician Med. Fam. Can.* **2015**, *61*, 233–239.
136. Wang, R.; Tian, H.; Guo, D.; Tian, Q.; Yao, T.; Kong, X. Impacts of exercise intervention on various diseases in rats. *J. Sport Health Sci.* **2020**, *9*, 211–227. [[CrossRef](#)] [[PubMed](#)]
137. Asif, Y.; Wlodek, M.E.; Black, M.J.; Russell, A.P.; Soeding, P.F.; Wadley, G.D. Sustained cardiac programming by short-term juvenile exercise training in male rats. *J. Physiol.* **2018**, *596*, 163–180. [[CrossRef](#)] [[PubMed](#)]
138. Liao, P.-H.; Hsieh, D.J.-Y.; Kuo, C.-H.; Day, C.-H.; Shen, C.-Y.; Lai, C.-H.; Chen, R.-J.; Padma, V.V.; Kuo, W.-W.; Huang, C.-Y. Moderate exercise training attenuates aging-induced cardiac inflammation, hypertrophy and fibrosis injuries of rat hearts. *Oncotarget* **2015**, *6*, 35383–35394. [[CrossRef](#)]
139. Dworatzek, E.; Mahmoodzadeh, S.; Schubert, C.; Westphal, C.; Leber, J.; Kusch, A.; Kararigas, G.; Fliegner, D.; Moulin, M.; Ventura-Clapier, R.; et al. Sex differences in exercise-induced physiological myocardial hypertrophy are modulated by oestrogen receptor beta. *Cardiovasc. Res.* **2014**, *102*, 418–428. [[CrossRef](#)]
140. Verboven, M.; Cuypers, A.; Deluyker, D.; Lambrichts, I.; Eijnde, B.O.; Hansen, D.; Bitto, V. High intensity training improves cardiac function in healthy rats. *Sci. Rep.* **2019**, *9*, 1–8. [[CrossRef](#)]
141. Zile, M.R.; Baicu, C.F.; Ikonomidis, J.; Stroud, R.E.; Nietert, P.J.; Bradshaw, A.D.; Slater, R.; Palmer, B.M.; Van Buren, P.; Meyer, M.; et al. Myocardial Stiffness in Patients with Heart Failure and a Preserved Ejection Fraction: Contributions of Collagen and Titin. *Circulation* **2015**, *131*, 1247–1259. [[CrossRef](#)]
142. Hafstad, A.D.; Lund, J.; Hadler-Olsen, E.; Höper, A.C.; Larsen, T.S.; Aasum, E. High- and moderate-intensity training normalizes ventricular function and mechanoenergetics in mice with diet-induced obesity. *Diabetes* **2013**, *62*, 2287–2294. [[CrossRef](#)] [[PubMed](#)]
143. Timothy, J.W.D.; Ben, C.; Sorcha, C.; de Marvao, A.; Roddy, W.; James, S.W.; Stuart, A.C.; O'Regan, P.D. Moderate Physical Activity in Healthy Adults Is Associated With Cardiac Remodeling. *Circ. Cardiovasc. Imaging* **2016**, *9*, e004712.
144. Potoră, C.S.; Tache, S.; Albu, A.; Nagy, A.L. Effects of moderate exercise and a multiple vitamin and mineral complex on the arterial wall. *Romanian J. Morphol. Embryol. Rev. Roum. Morphol. Embryol.* **2018**, *59*, 249–256.
145. Albarwani, S.; Al-Siyabi, S.; Baomar, H.; Hassan, M.O. Exercise training attenuates ageing-induced BKCa channel downregulation in rat coronary arteries. *Exp. Physiol.* **2010**, *95*, 746–755. [[CrossRef](#)] [[PubMed](#)]
146. Shi, L.; Liu, B.; Li, N.; Xue, Z.; Liu, X. Aerobic exercise increases BK(Ca) channel contribution to regulation of mesenteric arterial tone by upregulating β 1-subunit. *Exp. Physiol.* **2013**, *98*, 326–336. [[CrossRef](#)]
147. Zhao, H.-C.; Wang, F. Exercise training changes the gating properties of large-conductance Ca²⁺-activated K⁺ channels in rat thoracic aorta smooth muscle cells. *J. Biomech.* **2010**, *43*, 263–267. [[CrossRef](#)]
148. Zhang, Y.; Chen, Y.; Zhang, L.; Lu, N.; Shi, L. Aerobic exercise of low to moderate intensity corrects unequal changes in BK(Ca) subunit expression in the mesenteric arteries of spontaneously hypertensive rats. *Physiol. Res.* **2017**, *66*, 219–233. [[CrossRef](#)]
149. Shenouda, N.; Gillen, J.B.; Gibala, M.J.; MacDonald, M.J. Changes in brachial artery endothelial function and resting diameter with moderate-intensity continuous but not sprint interval training in sedentary men. *J. Appl. Physiol.* **2017**, *123*, 773–780. [[CrossRef](#)]
150. Sawyer, B.J.; Tucker, W.J.; Bhammar, D.M.; Ryder, J.R.; Sweazea, K.L.; Gaesser, G.A. Effects of high-intensity interval training and moderate-intensity continuous training on endothelial function and cardiometabolic risk markers in obese adults. *J. Appl. Physiol. Bethesda Md.* **2016**, *121*, 279–288. [[CrossRef](#)] [[PubMed](#)]
151. Rakobowchuk, M.; Tanguay, S.; Burgomaster, K.A.; Howarth, K.R.; Gibala, M.J.; MacDonald, M.J. Sprint interval and traditional endurance training induce similar improvements in peripheral arterial stiffness and flow-mediated dilation in healthy humans. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* **2008**, *295*, R236–R242. [[CrossRef](#)]

152. Goto, C.; Higashi, Y.; Kimura, M.; Noma, K.; Hara, K.; Nakagawa, K.; Kawamura, M.; Chayama, K.; Yoshizumi, M.; Nara, I. Effect of Different Intensities of Exercise on Endothelium-Dependent Vasodilation in Humans: Role of Endothelium-Dependent Nitric Oxide and Oxidative Stress. *Circulation* **2003**, *108*, 530–535. [CrossRef]
153. Sugawara, J.; Inoue, H.; Hayashi, K.; Yokoi, T.; Kono, I. Effect of low-intensity aerobic exercise training on arterial compliance in postmenopausal women. *Hypertens. Res. Off. J. Jpn. Soc. Hypertens.* **2004**, *27*, 897–901. [CrossRef] [PubMed]
154. WHO | What is Moderate-intensity and Vigorous-intensity Physical Activity? Available online: https://www.who.int/dietphysicalactivity/physical_activity_intensity/en/ (accessed on 21 October 2020).
155. *Physical Activity Guidelines for Americans*, 2nd ed.; Department of Health and Human Services: Washington, DC, USA, 2018.
156. de Oliveira, S.G.; dos Santos, N.V.; de Oliveira, F.S.R.; Souza-Mello, V.; da-Silva, B.S. High-intensity interval training has beneficial effects on cardiac remodeling through local renin-angiotensin system modulation in mice fed high-fat or high-fructose diets. *Life Sci.* **2017**, *189*, 8–17. [CrossRef] [PubMed]
157. Ross, L.M.; Porter, R.R.; Durstine, J.L. High-intensity interval training (HIIT) for patients with chronic diseases. *J. Sport Health Sci.* **2016**, *5*, 139–144. [CrossRef] [PubMed]
158. Green, D.J.; Hopman, M.T.E.; Padilla, J.; Laughlin, M.H.; Thijssen, D.H.J. Vascular Adaptation to Exercise in Humans: Role of Hemodynamic Stimuli. *Physiol. Rev.* **2017**, *97*, 495–528. [CrossRef]
159. Kong, Z.; Fan, X.; Sun, S.; Song, L.; Shi, Q.; Nie, J. Comparison of High-Intensity Interval Training and Moderate-to-Vigorous Continuous Training for Cardiometabolic Health and Exercise Enjoyment in Obese Young Women: A Randomized Controlled Trial. *PLoS ONE* **2016**, *11*, e0158589. [CrossRef]
160. Skutnik, B.C.; Smith, J.R.; Johnson, A.M.; Kurti, S.P.; Harms, C.A. The Effect of Low Volume Interval Training on Resting Blood Pressure in Pre-hypertensive Subjects: A Preliminary Study. *Phys. Sportsmed.* **2016**, *44*, 177–183. [CrossRef] [PubMed]
161. Brown, M.B.; Neves, E.; Long, G.; Graber, J.; Gladish, B.; Wiseman, A.; Owens, M.; Fisher, A.J.; Presson, R.G.; Petrache, I.; et al. High-intensity interval training, but not continuous training, reverses right ventricular hypertrophy and dysfunction in a rat model of pulmonary hypertension. *Am. J. Physiol.-Regul. Integr. Comp. Physiol.* **2017**, *312*, R197–R210. [CrossRef]
162. Cassidy, S.; Thoma, C.; Hallsworth, K.; Parikh, J.; Hollingsworth, K.G.; Taylor, R.; Jakovljevic, D.G.; Trenell, M.I. High intensity intermittent exercise improves cardiac structure and function and reduces liver fat in patients with type 2 diabetes: A randomised controlled trial. *Diabetologia* **2016**, *59*, 56–66. [CrossRef]
163. Novoa, U.; Arauna, D.; Moran, M.; Nuñez, M.; Zagmutt, S.; Saldivia, S.; Valdes, C.; Villaseñor, J.; Zambrano, C.G.; Gonzalez, D.R. High-Intensity Exercise Reduces Cardiac Fibrosis and Hypertrophy but Does Not Restore the Nitroso-Redox Imbalance in Diabetic Cardiomyopathy. *Oxid. Med. Cell. Longev.* **2017**, *2017*, 7921363. [CrossRef]
164. Oláh, A.; Mátyás, C.; Kellermayer, D.; Ruppert, M.; Barta, B.A.; Sayour, A.A.; Török, M.; Koncsos, G.; Gircz, Z.; Ferdinandy, P.; et al. Sex Differences in Morphological and Functional Aspects of Exercise-Induced Cardiac Hypertrophy in a Rat Model. *Front. Physiol.* **2019**, *10*, 889. [CrossRef] [PubMed]
165. Gomes, E.R.M.; Lara, A.A.; Almeida, P.W.M.; Guimarães, D.; Resende, R.R.; Campagnole-Santos, M.J.; Bader, M.; Santos, R.A.S.; Guatimosim, S. Angiotensin-(1-7) Prevents Cardiomyocyte Pathological Remodeling Through a Nitric Oxide/Guanosine 3',5'-Cyclic Monophosphate-Dependent Pathway. *Hypertension* **2010**, *55*, 153–160. [CrossRef]
166. Santos, R.A.S.; e Silva, A.C.S.; Maric, C.; Silva, D.M.R.; Machado, R.P.; de Buhr, I.; Heringer-Walther, S.; Pinheiro, S.V.B.; Lopes, M.T.; Bader, M.; et al. Angiotensin-(1-7) is an endogenous ligand for the G protein-coupled receptor Mas. *Proc. Natl. Acad. Sci. USA* **2003**, *100*, 8258–8263. [CrossRef] [PubMed]
167. Sukumaran, V.; Veeraveedu, P.T.; Gurusamy, N.; Yamaguchi, K.; Lakshmanan, A.P.; Ma, M.; Suzuki, K.; Kodama, M.; Watanabe, K. Cardioprotective effects of telmisartan against heart failure in rats induced by experimental autoimmune myocarditis through the modulation of angiotensin-converting enzyme-2/angiotensin 1-7/mas receptor axis. *Int. J. Biol. Sci.* **2011**, *7*, 1077–1092. [CrossRef] [PubMed]
168. Sukumaran, V.; Veeraveedu, P.T.; Gurusamy, N.; Lakshmanan, A.P.; Yamaguchi, K.; Ma, M.; Suzuki, K.; Kodama, M.; Watanabe, K. Telmisartan acts through the modulation of ACE-2/ANG 1-7/mas receptor in rats with dilated cardiomyopathy induced by experimental autoimmune myocarditis. *Life Sci.* **2012**, *90*, 289–300. [CrossRef] [PubMed]
169. Schinzari, F.; Veneziani, A.; Mores, N.; Barini, A.; Di Daniele, N.; Cardillo, C.; Tesauro, M. Beneficial Effects of Apelin on Vascular Function in Patients With Central Obesity. *Hypertension* **2017**, *69*, 942–949. [CrossRef]
170. Rahimi, M.; Shekarforoush, S.; Asgari, A.R.; Khoshbaten, A.; Rajabi, H.; Bazgir, B.; Mohammadi, M.T.; Sobhani, V.; Shakibae, A. The effect of high intensity interval training on cardioprotection against ischemia-reperfusion injury in wistar rats. *EXCLI J.* **2015**, *14*, 237–246.
171. Heiskanen, M.A.; Leskinen, T.; Heinonen, I.H.A.; Löyttyniemi, E.; Eskelinen, J.-J.; Virtanen, K.; Hannukainen, J.C.; Kalliokoski, K.K. Right ventricular metabolic adaptations to high-intensity interval and moderate-intensity continuous training in healthy middle-aged men. *Am. J. Physiol.-Heart Circ. Physiol.* **2016**, *311*, H667–H675. [CrossRef] [PubMed]
172. Grace, F.; Herbert, P.; Elliott, A.D.; Richards, J.; Beaumont, A.; Sculthorpe, N.F. High intensity interval training (HIIT) improves resting blood pressure, metabolic (MET) capacity and heart rate reserve without compromising cardiac function in sedentary aging men. *Exp. Gerontol.* **2018**, *109*, 75–81. [CrossRef]
173. Ito, S. High-intensity interval training for health benefits and care of cardiac diseases—The key to an efficient exercise protocol. *World J. Cardiol.* **2019**, *11*, 171–188. [CrossRef]

174. Stewart, G.M.; Chan, J.; Yamada, A.; Kavanagh, J.J.; Haseler, L.J.; Shiino, K.; Sabapathy, S. Impact of high-intensity endurance exercise on regional left and right ventricular myocardial mechanics. *Eur. Heart J. Cardiovasc. Imaging* **2016**, *18*, 688–696. [[CrossRef](#)]
175. Cote, A.T.; Phillips, A.A.; Foulds, H.J.; Charlesworth, S.A.; Bredin, S.S.D.; Burr, J.F.; Koehle, M.S.; Warburton, D.E.R. Sex Differences in Cardiac Function After Prolonged Strenuous Exercise. *Clin. J. Sport Med.* **2015**, *25*, 276–283. [[CrossRef](#)]
176. Scott, J.M.; Esch, B.T.A.; Haykowsky, M.J.; Isserow, S.; Koehle, M.S.; Hughes, B.G.; Zbogar, D.; Bredin, S.S.D.; McKenzie, D.C.; Warburton, D.E.R. Sex differences in left ventricular function and β -receptor responsiveness following prolonged strenuous exercise. *J. Appl. Physiol.* **2007**, *102*, 681–687. [[CrossRef](#)]
177. Batacan, R.B.; Duncan, M.J.; Dalbo, V.J.; Buitrago, G.L.; Fenning, A.S. Effect of different intensities of physical activity on cardiometabolic markers and vascular and cardiac function in adult rats fed with a high-fat high-carbohydrate diet. *J. Sport Health Sci.* **2018**, *7*, 109–119. [[CrossRef](#)] [[PubMed](#)]
178. Klonizakis, M.; Moss, J.; Gilbert, S.; Broom, D.; Foster, J.; Tew, G.A. Low-volume high-intensity interval training rapidly improves cardiopulmonary function in postmenopausal women. *Menopause* **2014**, *21*, 1099–1105. [[CrossRef](#)] [[PubMed](#)]
179. Rognmo, Ø.; Hetland, E.; Helgerud, J.; Hoff, J.; Slørdahl, S.A. High intensity aerobic interval exercise is superior to moderate intensity exercise for increasing aerobic capacity in patients with coronary artery disease. *Eur. J. Cardiovasc. Prev. Rehabil.* **2004**, *11*, 216–222. [[CrossRef](#)] [[PubMed](#)]
180. Freyssin, C.; Verkindt, C.; Prieur, F.; Benaich, P.; Maunier, S.; Blanc, P. Cardiac Rehabilitation in Chronic Heart Failure: Effect of an 8-Week, High-Intensity Interval Training Versus Continuous Training. *Arch. Phys. Med. Rehabil.* **2012**, *93*, 1359–1364. [[CrossRef](#)] [[PubMed](#)]
181. Fang, Y.; Ying, W.; Yu, C.; Daliao, X.; Lijun, S. Impact of moderate- and high-intensity exercise on the endothelial ultrastructure and function in mesenteric arteries from hypertensive rats. *Life Sci.* **2019**, *222*, 36–45.
182. Chen, Y.; Zhang, H.; Zhang, Y.; Lu, N.; Zhang, L.; Shi, L. Exercise intensity-dependent reverse and adverse remodeling of voltage-gated Ca^{2+} channels in mesenteric arteries from spontaneously hypertensive rats. *Hypertens. Res.* **2015**, *38*, 656–665. [[CrossRef](#)]
183. Beckers, P.J.; Denollet, J.; Possemiers, N.M.; Wuyts, F.L.; Vrints, C.J.; Conraads, V.M. Combined endurance-resistance training vs. endurance training in patients with chronic heart failure: A prospective randomized study. *Eur. Heart J.* **2008**, *29*, 1858–1866. [[CrossRef](#)]
184. Dor-Haim, H.; Barak, S.; Horowitz, M.; Yaakobi, E.; Katzburg, S.; Swissa, M.; Lotan, C. Improvement in cardiac dysfunction with a novel circuit training method combining simultaneous aerobic-resistance exercises. A randomized trial. *PLoS ONE* **2018**, *13*, e0188551.
185. Schroeder, E.C.; Franke, W.D.; Sharp, R.L.; Lee, D. Comparative effectiveness of aerobic, resistance, and combined training on cardiovascular disease risk factors: A randomized controlled trial. *PLoS ONE* **2019**, *14*, e0210292. [[CrossRef](#)]
186. Lippincott Williams & Wilkins. *ACSM's Guidelines for Exercise Testing and Prescription*, 9th ed.; American College of Sports Medicine: Baltimore, MD, USA, 2013; ISBN 978-1-60913-605-5.
187. Patel, H.; Alkhwam, H.; Madanieh, R.; Shah, N.; Kosmas, C.E.; Vittorio, T.J. Aerobic vs anaerobic exercise training effects on the cardiovascular system. *World J. Cardiol.* **2017**, *9*, 134–138. [[CrossRef](#)]
188. Stojiljković, N.; Ignjatović, A.; Savić, Z.; Marković, Ž.; Milanović, S. History of Resistance Training. *Act. Phys. Educ. Sport* **2013**, *3*, 135–138.
189. Shaw, B.S.; Shaw, I.; Brown, G.A. Resistance exercise is medicine: Strength training in health promotion and rehabilitation. *Int. J. Ther. Rehabil.* **2015**, *22*, 385–389. [[CrossRef](#)]
190. Sanches, I.C.; Buzin, M.; Conti, F.F.; da Dias, D.S.; dos Santos, C.P.; Sirvente, R.; Salemi, V.M.C.; Llesuy, S.; Irigoyen, M.-C.; De Angelis, K. Combined aerobic and resistance exercise training attenuates cardiac dysfunctions in a model of diabetes and menopause. *PLoS ONE* **2018**, *13*, e0202731.
191. Shimojo, G.L.; da Silva Dias, D.; Malfitano, C.; Sanches, I.C.; Llesuy, S.; Ulloa, L.; Irigoyen, M.-C.; De Angelis, K. Combined Aerobic and Resistance Exercise Training Improve Hypertension Associated With Menopause. *Front. Physiol.* **2018**, *9*, 1471. [[CrossRef](#)] [[PubMed](#)]
192. Chrysohoou, C.; Angelis, A.; Tsitsinakis, G.; Spetsioti, S.; Nasis, I.; Tsiachris, D.; Rapakoulis, P.; Pitsavos, C.; Koulouris, N.G.; Vogiatzis, I.; et al. Cardiovascular effects of high-intensity interval aerobic training combined with strength exercise in patients with chronic heart failure. A randomized phase III clinical trial. *Int. J. Cardiol.* **2015**, *179*, 269–274. [[CrossRef](#)] [[PubMed](#)]
193. Haykowsky, M.J.; Liang, Y.; Pechter, D.; Jones, L.W.; McAlister, F.A.; Clark, A.M. A Meta-Analysis of the Effect of Exercise Training on Left Ventricular Remodeling in Heart Failure Patients: The Benefit Depends on the Type of Training Performed. *J. Am. Coll. Cardiol.* **2007**, *49*, 2329–2336. [[CrossRef](#)]
194. Ahn, N.; Kim, K. Effects of Aerobic and Resistance Exercise on Myokines in High Fat Diet-Induced Middle-Aged Obese Rats. *Int. J. Environ. Res. Public Health* **2020**, *17*, 2685. [[CrossRef](#)] [[PubMed](#)]
195. Johansson, M.; Gao, S.A.; Friberg, P.; Annerstedt, M.; Carlström, J.; Ivarsson, T.; Jensen, G.; Ljungman, S.; Mathillas, O.; Nielsen, F.-D.; et al. Baroreflex effectiveness index and baroreflex sensitivity predict all-cause mortality and sudden death in hypertensive patients with chronic renal failure. *J. Hypertens.* **2007**, *25*, 163–168. [[CrossRef](#)]
196. Masson, G.S.; Costa, T.S.R.; Yshii, L.; Fernandes, D.C.; Soares, P.P.S.; Laurindo, F.R.; Scavone, C.; Michelini, L.C. Time-Dependent Effects of Training on Cardiovascular Control in Spontaneously Hypertensive Rats: Role for Brain Oxidative Stress and Inflammation and Baroreflex Sensitivity. *PLoS ONE* **2014**, *9*, e94927. [[CrossRef](#)]

197. Son, W.-M.; Sung, K.-D.; Cho, J.-M.; Park, S.-Y. Combined exercise reduces arterial stiffness, blood pressure, and blood markers for cardiovascular risk in postmenopausal women with hypertension. *Menopause* **2017**, *24*, 262–268. [[CrossRef](#)]
198. Maeda, S.; Zempo-Miyaki, A.; Sasai, H.; Tsujimoto, T.; So, R.; Tanaka, K. Lifestyle modification decreases arterial stiffness in overweight and obese men: Dietary modification vs. exercise training. *Int. J. Sport Nutr. Exerc. Metab.* **2015**, *25*, 69–77. [[CrossRef](#)] [[PubMed](#)]
199. Masroor, S.; Bhati, P.; Verma, S.; Khan, M.; Hussain, M.E. Heart Rate Variability following Combined Aerobic and Resistance Training in Sedentary Hypertensive Women: A Randomised Control Trial. *Indian Heart J.* **2018**, *70*, S28–S35. [[CrossRef](#)]
200. Figueroa, A.; Park, S.Y.; Seo, D.Y.; Sanchez-Gonzalez, M.A.; Baek, Y.H. Combined resistance and endurance exercise training improves arterial stiffness, blood pressure, and muscle strength in postmenopausal women. *Menopause* **2011**, *18*, 980–984. [[CrossRef](#)] [[PubMed](#)]
201. Miyachi, M.; Kawano, H.; Sugawara, J.; Takahashi, K.; Hayashi, K.; Yamazaki, K.; Tabata, I.; Tanaka, H. Unfavorable Effects of Resistance Training on Central Arterial Compliance: A Randomized Intervention Study. *Circulation* **2004**, *110*, 2858–2863. [[CrossRef](#)] [[PubMed](#)]
202. Miyachi, M.; Donato, A.J.; Yamamoto, K.; Takahashi, K.; Gates, P.E.; Moreau, K.L.; Tanaka, H. Greater Age-Related Reductions in Central Arterial Compliance in Resistance-Trained Men. *Hypertension* **2003**, *41*, 130–135. [[CrossRef](#)] [[PubMed](#)]
203. Kawano, H.; Tanaka, H.; Miyachi, M. Resistance training and arterial compliance: Keeping the benefits while minimizing the stiffening. *J. Hypertens.* **2006**, *24*, 1753–1759. [[CrossRef](#)]
204. Montero, D.; Vinet, A.; Roberts, C.K. Effect of combined aerobic and resistance training versus aerobic training on arterial stiffness. *Int. J. Cardiol.* **2015**, *178*, 69–76. [[CrossRef](#)]
205. Lima, L.; Bonardi, J.; Campos, G.; Bertani, R.; Scher, L.; Moriguti, J.; Ferriolli, E.; Lima, N. Combined aerobic and resistance training: Are there additional benefits for older hypertensive adults? *Clinics* **2017**, *72*, 363–369. [[CrossRef](#)]
206. Shiotsu, Y.; Watanabe, Y.; Tujii, S.; Yanagita, M. Effect of exercise order of combined aerobic and resistance training on arterial stiffness in older men. *Exp. Gerontol.* **2018**, *111*, 27–34. [[CrossRef](#)] [[PubMed](#)]
207. Okamoto, T.; Masuhara, M.; Ikuta, K. Combined aerobic and resistance training and vascular function: Effect of aerobic exercise before and after resistance training. *J. Appl. Physiol.* **2007**, *103*, 1655–1661. [[CrossRef](#)] [[PubMed](#)]
208. Li, Y.; Hanssen, H.; Cordes, M.; Rossmeissl, A.; Endes, S.; Schmidt-Trucksäss, A. Aerobic, resistance and combined exercise training on arterial stiffness in normotensive and hypertensive adults: A review. *Eur. J. Sport Sci.* **2015**, *15*, 443–457. [[CrossRef](#)] [[PubMed](#)]