Original article

High-intensity Interval Training Dosage for Heart Failure and Coronary Artery Disease Cardiac Rehabilitation. A Systematic Review and Meta-analysis



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ABSTRACT

Introduction and objectives: High-interval intensity training (HIT) has been suggested to improve peak VO₂ in cardiac rehabilitation programs. However, the optimal HIT protocol is unknown. The objective of this study was to identify the most effective doses of HIT to optimize peak VO₂ in coronary artery disease (CAD) and heart failure (HF) patients.

Methods: A search was conducted in 6 databases (MEDLINE, Web of Science, LILACS, CINAHL, Academic Search Complete, and SportDiscus). Studies using a HIT protocol in CAD or HF patients and measuring peak VO₂ were included. The PEDro Scale and Cochrane Collaboration tools were used.

Results: Analyses reported significant improvements in peak VO₂ after HIT in both diseases (P = .000001), with a higher increase in HF patients (P = .03). Nevertheless, in HF patients, there were no improvements when the intensity recovery was $\le 40\%$ of peak VO₂ (P = .19) and the frequency of training was ≤ 2 d/wk (P = .07). There were significant differences regarding duration in CAD patients, with greater improvements in peak VO₂ when the duration was < 12 weeks (P = .05). In HF, programs lasting < 12 weeks did not significantly improve peak VO₂ (P = .1).

Conclusions: The HIT is an effective method for improving peak VO₂ in HF and CAD, with a significantly greater increase in HF patients. The recovery intervals should be active and be between 40% and 60% of peak VO₂ in HF patients. Training frequency should be ≥ 2 d/wk for CAD patients and ≥ 3 d/wk for HF patients.

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Dosis de ejercicio interválico de alta intensidad en la rehabilitación cardiaca de la insuficiencia cardiaca y la enfermedad arterial coronaria: revisión sistemática y metanálisis

RESUMEN

Introducción y objetivos: Se ha propuesto el ejercicio interválico de alta intensidad (EIAI) en programas de rehabilitación cardiaca para mejorar el VO_{2máx}. Sin embargo, no se conoce cuál es el mejor protocolo EIAI. El objetivo es encontrar la mejor dosis de EIAI para optimizar el VO_{2máx} de pacientes con enfermedad arterial coronaria (EAC) e insuficiencia cardiaca (IC).

Método: Se llevó a cabo una búsqueda en 6 bases de datos (MEDLINE, *Web of Science*, LILACS, CINAHL, Academic Search Complete y SportDiscus). Se incluyeron los estudios que usaban el protocolo EIAI y midieron el VO_{2máx} de pacientes con EAC e IC. Se utilizó la escala PEDro y las herramientas de la Colaboración Cochrane.

Resultados: El análisis mostró mejoras significativas en el VO_{2máx} tras el EIAI en ambas enfermedades (p = 0,000001), con mayor incremento en los pacientes con IC (p = 0,03). Sin embargo, en estos no hubo mejora si la intensidad de recuperación era $\leq 40\%$ del VO_{2máx} (p = 0,19) o la frecuencia de entrenamiento era ≤ 2 días/semana (p = 0,07). Hubo diferencias significativas según la duración entre los pacientes con IC no mostraron resultados superiores cuando era < 12 semanas (p = 0,05). Los pacientes con IC no mostraron mejoras significativas en el VO_{2máx} cuando la duración era < 12 semanas (p = 0,1).

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Conclusiones: El EIAI es un método efectivo para mejorar el VO_{2máx} de los pacientes con IC o EAC, con mayor diferencia significativa en los pacientes con IC. Los intervalos de recuperación de los pacientes con IC deben ser activos y estar en un 40-60% del VO_{2máx}. La frecuencia de entrenamiento debería ser ≥ 2 días/semana en la EAC y ≥ 3 días/semana en la IC.

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Abbreviations

CAD: coronary artery disease HF: heart failure HIT: high-intensity interval training Peak VO₂: peak oxygen uptake

INTRODUCTION

Coronary artery disease (CAD) is one of the most common causes of death worldwide, affecting 17.5 million people each year.¹ Ischemic disease and chronic heart failure (HF) are lethal, causing 8.76 million deaths worldwide,² resulting in higher health care system spending.³ These CAD reduce either exercise tolerance or peak oxygen uptake (peak VO₂), culminating in heart, respiratory, and endothelial dysfunction.⁴ In addition, HF is the final condition of all cardiovascular diseases, as it affects both cardiac function and cardiovascular circulation.⁵ For this reason, they are defined as an incapacity syndrome, which creates morbidity and whose symptoms are linked to fatigue, dyspnea, and exercise intolerance.⁴

Currently, it has been widely proven that one of the benefits of aerobic exercise is an increase in peak VO₂,⁶⁻⁹ which is one of the most important survival indicators for people with CAD^{5,10-13} and HF.⁵ Continuous aerobic training has been studied for cardiac rehabilitation programs. It consists of performing exercise for a long period of time under moderate intensity and nonvariable aerobic activity (60%-80% of peak $VO_2^{5,9,11,12,14}$). However, at present, high-intensity interval training (HIT) protocols are generating better results in peak VO₂ in less time.^{9,13–16} They consist of intermittent short high-intensity work periods (85%-100% of peak VO₂) and relative rest periods.^{10,15} A typical HIT session consists of a 10-minute warm-up at 50% to 70% of peak VO₂ followed by a set of four 3- to 4-minute bouts^{5,10-13,15-23} (HIT protocol with long work interval) or a set of ten 30- to 60-second bouts^{6-9,24} (HIY protocol with short work interval) at 85% to 95% of peak VO₂ interspersed by active pauses at 50% to 70% of peak VO₂. It terminates with a cool-down period at 50% to 70% of peak VO₂² (Figure 1). However, although HIT is known to produce the greatest effects on peak VO₂ in CAD and HF patients,^{5,13,15,16,21} there have been many differences in the exercise protocols proposed in this topic affecting the frequency (2-5 d/wk),^{8,11,14,24} volume (30-60 min/sessions),^{15,24,26} intensity of recovery (0%-70% peak VO_2),^{6,9,17,21} number of sessions (> 100 sessions),¹³ duration of training (4-50 weeks),^{8,13,16,18} and even in the characteristics of the patients.^{11,17,21} Recognizing the influence of these variables regarding peak VO₂ could help to optimize cardiac rehabilitation programs for HF and CAD patients.

Moreover, other studies have focused on identifying which kind of HIT intervention is more effective for improving the functional capacity of HF and CAD patients, as the rehabilitation process and the disabilities produced by the 2 diseases differ.^{10,27} However, which HIT protocol training is better for one disease or the other, or the best dosage of this type of exercise to improve the cardiac rehabilitation program of each disease is currently not known. Therefore, the aim of the present study was to conduct a metaanalysis to a) identify the best doses of HIT to optimize HIT training for HF and CAD patients, and b) determine how the HIT protocol affects HF and CAD patients in terms of their peak VO₂.

METHODS

Study Design

The systematic review and meta-analysis were undertaken in accordance with the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) statement.²⁸ The search was carried out by 2 of the authors (I. Ballesta García and J.A. Rubio Arias), and all the authors reached an agreement regarding methods, article inclusion, and statistical analysis. The articles were organized in order to delete duplicates. Full-text articles were retrieved and evaluated based on the proposed selection criteria. Variables related to the training method such as frequency, total duration of the session, exercise duration, and recovery intervals, number of total sessions, percentage of intensity of each interval, as well as the age, body mass index, sex and number of patients in each study were extracted to an excel workbook.

Search Strategy

Electronic database searches were performed using MEDLINE, Web of Science, LILACS, CINAHL, Academic Search Complete and SportDiscus. The search terms chosen were: ("Heart disease" OR "Heart Failure" OR "Coronary Artery Disease" OR CAD) AND ("High-intensity" OR "High intensity" OR HIT OR "Interval training"). The search results and final study selection are shown in Figure 2.

Inclusion Criteria

Inclusion criteria were determined by the authors. We included randomized controlled trials, written in English or Spanish, published from January 2004 to March 2017, using a HIT protocol in patients with CAD or HF, measuring peak VO₂. We excluded studies based on aquatic HIT programs, combining HIT with strength training, home-based HIT, not considering CAD and HF results of peak VO₂ independently, using people with transplants, grafts or with valve disease, and testing with food supplements, nutritional or pharmacological aids.

Statistical Analysis

The meta-analysis and statistical analysis were developed using Review Manager software (RevMan 5.2; Cochrane Collaboration, Oxford, United Kingdom)²⁹ and Comprehensive Meta-analysis software (Version 2; Biostat, Englewood, New Jersey, United States). A random effects meta-analysis was conducted to determine which disease most benefited from HIT and if there were differences between HIT protocols for each condition. The work-rest ratio, the work interval duration and intensity, and the



Figure 1. Schematic illustration for HIT_S and HIT_L. HIT, high-intensity interval training groups; HIT_L, protocol of HIT with long work intervals; HIT_S, protocol of HIT with short work intervals.



Figure 2. Systematic review and meta-analysis flow diagram. WOS, Web of Science.

General Characteristics of the Studies Included in the Meta-analysis

Research studies	Study location, country	Disease	n_HIT	Men, %	Age, y	BMI, cm/kg ²
Benda et al. ²⁴	Nijmegen, Netherlands	HF	10	100	63.0 ± 8	28.1 ± 7.5
Cardozo et al. ¹¹	Rio de Janerio, Brasil	CAD	23	63	56.0 ± 12	27.5 ± 5.9
Chrysohoou et al. ⁶	Rio de Janerio, Brasil	HF	33	88	63.0 ± 9	28.8 ± 4.2
Conraads et al. ¹⁸	Athens, Greece	CAD	100	85	57.0 ± 8.8	28 ± 4.4
Currie et al.9	Hamilton, Canada	CAD	11	91	67.2 ± 6	27.9 ± 4.9
Dimopoulos et al. ²⁷	Athens, Greece	HF	10	90	59.2 ± 12.2	26.5 ± 4.1
Freyssin et al. ⁸	Reunion island, France	HF	13	50	54.0 ± 9	24.8 ± 4
Fu et al. ¹⁶	Keelung, Taiwan	HF	14	61	67.5 ± 1.8	
Huang et al. ¹⁹	Taoyuan, Taiwan	HF	33	78	60.0 ± 3	
Isaksen et al. ¹⁷	Stavanger, Norway	HF	24	88	65.0 ± 9	27.8 ± 4
Kim et al. ¹⁵	Seoul, Korea	CAD	14	86	57.0 ± 11.5	24.2 ± 2.9
Koufaki et al. ³⁰	Staffordshire, England	HF	16	87	59.8 ± 7.4	28.9 ± 4.7
Madssen et al. ¹³	Trondheim, Norway	CAD	24	75	64.4 [47-78]	28 ± 3.9
Moholdt et al. ³¹	Trondheim, Norway	CAD	28	86	60.2 ± 6.9	26 ± 6.2
Moholdt et al. ²¹	Trondheim, Norway	CAD	11	83	57.7 ± 9.3	26.6 ± 3
Roditis et al. ⁷	Athens, Greece	HF	11	90	63.0 ± 2	25.9 ± 2.8
Rognmo et al. ³²	Trondheim, Norway	CAD	8	75	$\textbf{62.9} \pm \textbf{11.2}$	26.7 ± 4.1
Smart et al. ³³	Athens, Greece	HF	10	100	59.1 ± 11	28.9 ± 6.1
Warburton et al. ³⁴	Vancouver, Canada	CAD	7	100	55.0 ± 7	

BMI, body mass index; CAD, coronary artery disease; HF, heart failure; n_HIT number of participants in high-intensity interval training groups. Data are presented as the mean, mean \pm standard deviation, range or No.

protocol frequency variables were taken into account to determine changes in each variable.

Due to the heterogeneity of the protocol (Table 1), mean differences were used, dividing the mean values between 2 different groups. The differences in means were grouped using the random effects model. Heterogeneity between studies was analyzed using l^2 statistics. The dichotomous and continuous variables of the studies were compared with extracted potential peak VO₂ moderator factors. The medians of continuous variables were used to group the studies. Afterwards, HIT differences between pre- and post-values were expressed and analyzed as potential moderator variable changes. The publication bias for the different conditions analyzed (pre-vs-post) was assessed by examining the asymmetry of a funnel plot using Egger's test, and P < .05 was considered to be statistically significant. We assessed the methodological quality of the studies by using the PEDro Scale. The risk of bias was assessed using the modified Cochrane Collaboration tools. Bias was assessed as a judgment (high, low, or unclear) for individual elements from 5 domains: selection, performance, attrition, reporting and any other bias (criteria inclusion of patients in the studies and the country in which the study was conducted).

RESULTS

According to our inclusion criteria, 19 studies were included in this meta-analysis, 10 in HF and 9 in CAD. The Egger test provided statistical evidence of funnel plot symmetry (Figure 3), suggesting the absence of a significant publication bias.

Risk-of-bias assessment is shown in Figure 4. It was high in almost all studies due to lack of blinding of participants and personnel. However, this issue could not be omitted due to the peculiarity of the intervention (exercise vs no exercise) and should be considered in perspective.

The main characteristics of the studies and of training interventions are described in Table 1 and Table 2, respectively.



Figure 3. Funnel plot of comparison: CAD vs HF. CAD, coronary artery disease; HF, heart failure; MD, mean difference; SE, standard error.

The meta-analyzed effects of HIT found in both diseases were beneficial (P < .000001) for peak VO₂ (3.98; 95%CI, 2.94-5.02 mL/ kg/min for CAD patients and 2.55; 95%CI, 1.73-3.36 mL/kg/min for HF patients) (Figure 5). However, there were significant differences in peak VO₂ between the 2 diseases in favor of HF patients (P = .03) (Figure 5).

Following the moderating variables, the results showed statistical improvements in peak VO₂ in each subgroup analyzed for both diseases (P < .05) (Table 3 and Table 4). There were no statistically significant differences between subgroups according to population characteristics in CAD or HF (Table 3 and Table 4).

Regarding exercise characteristics, there were statistically significant improvements in peak VO₂ in each CAD patient subgroup analyzed (P < .05). Nevertheless, in HF patients, there were no improvements in peak VO₂ when the intensity of recovery was \leq 40% peak VO₂ (P = .19), when the type of recovery was passive (P = .09), and in the \leq 2 d/wk protocols (P = .07)^{9,24} (Table 5).



Figure 4. Assessment of risk of bias in included randomized controlled trials.

With regards to protocol duration, there were significant differences in CAD patients, who showed higher peak VO₂ improvements when the program was performed for < 12 weeks (P = .05) (Table 6). By contrast, there were no significant differences in HF patients when comparing a program \ge 12 weeks and a program \le 12 weeks (P = .96). However, a program lasting \ge 12 weeks did not produce significant improvements in peak VO₂ in HF patients (P = .1; $I^2 = 0$) (Table 5).

Likewise, there were no significant differences between HIT protocols with short exercise intervals and protocols with long exercise intervals in either of the 2 diseases (P = .87), although both of them showed significant improvements in peak VO₂ (P = < .000001). In contrast, there were significant differences in peak VO₂ between the 2 diseases in favor of HF patients (P = .03) (Figure 5).

DISCUSSION

This meta-analysis determined how different HIT protocols modify the peak VO₂ of HF and CAD patients, describing the most effective doses of HIT to optimize their training. The main finding was that HIT was more effective for improving the peak VO₂ of patients with HF than that of those with CAD. However, the maximum benefits in peak VO₂ were obtained between weeks 6 and 12 for both diseases. In addition, HF patients did not obtain significant peak VO₂ improvements when the HIT protocol was > 12 weeks. This result may be due to the fact that only 2 studies^{30,33} followed a protocol > 12 weeks while 8 studies^{6–8,16,17,19,24,27} followed protocols of \leq 12 weeks. In fact, no significant differences were found between the 2 subgroups (*P* = .96). Likewise, CAD patients achieved greater improvements in

Table 2

Characteristics of Aerobic Training Interventions in the Studies Included in the Meta-analysis

Studies	Disease	Туре	Frequency, wks	Session duration, min	Interval work, min	Interval recovery, min	Duration, wks	Number of sessions	Work intensity, % peak VO ₂ ª	R	Increase VO ₂ peak, % ^b
Benda et al. ²⁴	HF	HIT_S	2	30	1	2	12	24	90	0.5	6.8
Cardozo et al. ¹¹	CAD	HIT_L	3	28	4	3	16	48	90	1.33	18.44
Chrysohoou et al. ⁶	HF	HIT_S	3	45	0.5	0.5	12	36	100	1	31.25
Conraads et al. ¹⁸	CAD	HIT_L	3	28	4	3	12	36	90-95	1.33	21.7
Currie et al. ⁹	CAD	HIT_S	2	20	1	1	12	24	80-100	1	23.73
Dimopoulos et al. ²⁷	HF	HIT_S	3	36	0.5	0.5	12	36	100	1	7.79
Freyssin et al. ⁸	HF	HIT_S	5	69	0.5	1	8	40	80-95	0.5	27.1
Fu et al. ¹⁶	HF	HIT_L	3	30	3	3	12	36	80	1	22.5
Huang et al. ¹⁹	HF	HIT_L	3	42	3	3	12	36	80	1	13.41
Isaksen et al. ¹⁷	HF	HIT_L	3	28	4	3	12	36	85	1.33	5.74
Kim et al. ¹⁵	CAD	HIT_L	3	28	4	3	6	18	85-95	1.33	22.16
Koufaki et al. ³⁰	HF	HIT_S	3	-	0.5	1	24	72	100	0.5	15.68
Madssen et al. ¹³	CAD	HIT_L	3	28	4	3	52	156	85-95	1.33	3.22
Moholdt et al. ³¹	CAD	HIT_L	5	28	4	3	4	20	90	1.33	12.17
Moholdt et al. ²¹	CAD	HIT_L	3	28	4	3	12	36	85-95	1.33	14.55
Roditis et al. ⁷	HF	HIT_S	3	40	0.5	0.5	6	18	100	1	8.45
Rognmo et al. ³²	CAD	HIT_L	3	28	4	3	10	30	80-90	1.33	18.86
Smart et al. ³³	HF	HIT_S	3	60	1	1	48	16	70	0.5	20.49
Warburton et al. ³⁴	CAD	HIT_S	3	_	2	3	16	48	85-95	0.66	17.28

CAD, coronary artery disease; HF, heart failure; HIT, high-intensity interval training groups; HIT_L, protocol of HIT with long work interval; HIT_S, protocol of HIT with short work interval; peak VO₂, peak oxygen uptake; R: work/recovery ratio.

^a Percentage of peak VO2 of the interval work.

^b Percentage of increase of VO₂ peak posttraining.

		Postinte	rventi	on	Preinte	venti	on		MD	MD
	Study or subgroup	Mean	SD	Total	Mean	SD	Total	Weight I	IV, random, 95%CI	IV, random, 95%CI
A-	CAD									
	Cardozo et al ¹¹ HIT	24.4	5	23	20.6	5	23	4.7%	3.80 (0.91 to 6.69)	_
	Conraads et al.,18 HIT	28.6	6.9	100	23.5	5.7	100	8.9%	5.10 (3.35 to 6.85)	
	Currle et al., ⁹ HIT	24.5	4.5	11	19.8	3.7	11	3.6%	4.70 (1.26 to 8.14)	
	Kim et al., ¹⁵ HIT	35.5	7.7	14	29.2	5.5	14	1.9%	6.30 (1.34 to 11.26)	
	Madssen et al., ¹³ HIT	28.8	5.6	24	27.9	4.7	24	4.6%	0.90 (-2.02 to 3.82)	
	Moholdt et al., ³¹ HIT	30.4	5.5	28	27.1	4.5	28	5.4%	3.30 (0.67 to 5.93)	_
	Moholdt et al., ²¹ HIT	36.2	8.6	11	31.6	5.8	11	1.3%	4.60 (-1.53 to 10.73)	
	Rognmo et al., ²⁵ HIT	37.8	12.4	8	31.8	9.3	8	0.4%	6.00 (-4.74 to 16.74)	
	Warburton et al., ³⁴ HIT	19	4.3	7	16.2	3.5	7	2.6%	2.80 (-1.31 to 6.91)	
	Subtotal (95%CI)			226			226	33.4%	3.98 (2.94 to 5.02)	•
	Heterogeneity: Tau ² = 0.00; chi	-square = 7	7.60, df	= 8 (P	$= 0.47$; I^2	= 0%				
	Test for overall effect: Z = 7.51	(<i>P</i> < .0000)1)		,.					
В	HF									
	Benda et al ²⁴ HIT	20.4	4.3	10	19 1	4.1	10	3.2%	1 30 (-2 38 to 4 98)	_
	Chrysohoou et al. ⁶ HIT	20.4	5	33	16	6	33	5.3%	5 00 (2 34 to 7 66)	
	Dimonoulos et al. ²⁷ HIT	10.8	41	12	10.6	41	12	3.8%	0.20 (-3.08 to 3.48)	_
	Frevssin et al ⁸ HIT	13.6	3.2	14	10.7	2 99	14	6.5%	2.90(0.61 to 5.19)	
	Fuetal ¹⁶ HIT	19.6	12	14	16	1	15	15.4%	3.60(2.79 to 4.41)	-
	Huang et al. ¹⁹ HIT	18.6	0.9	33	16.4	0.6	33	18.1%	2 20 (1 83 to 2 57)	
	Isaksen et al ¹⁷ HIT	18.4	5.3	24	17.4	4.6	24	4.9%	1.00 (-1.81 to 3.81)	_
	Koukafi et al ³⁰ HIT	17.7	4.9	9	15.3	4.7	16	2.8%	2.40 (-1.54 to 6.34)	
	Roditis et al., ⁷ HIT	15.4	4.2	11	14.2	3.1	11	4.2%	1.20 (-1.88 to 4.28)	
	Smart et al ³³ HIT	14.7	4.5	10	12.2	5.4	10	2.4%	2.50 (-1.86 to 6.86)	
	Subtotal (95%CI)			170			178	66.6%	2.55 (1.73 to 3.36)	•
	Heterogeneity: Tau ² = 0.53; chi	-square = 1	17.05, 0	df = 9 (<i>F</i>	$P = .05); I^2$	= 47%				
	Test for overall effect: Z = 6.13	(P < .0000	1)	· ·	,.					
	Total (95%CI)			396			404	100.0%	2.98 (2.26 to 3.70)	•
	Heterogeneity: Tau ² = 0.71; chi	-square = 3	32.52, 0	df = 18 ($P = .02); I^2$	2 = 45%	6			—
	Test for overall effect: Z = 8.11	(P < .0000	1)		,,					-10 -5 0 5 10
	Test for subgroup differences: of	hi-square	, 4.54	df = 1 (P = .03; I	= 78.	0%			Favours (pre) Favours (post)

Figure 5. A: MD between post- and pre-HIT intervention peak VO₂ (mL/kg/min) for patients with CAD. B: MD between post- and pre-HIT intervention peak VO₂ (mL/kg/min) for patients with HF. Squares represent the MD for each trial. The diamond represents the pooled MD across trials. Weight determines how much each individual study contributes to the pooled estimate.³⁵ Total is the number of participants in HIT groups. 95%CI, confidence interval; CAD, coronary artery disease; HF, heart failure; HIT, high-intensity interval training groups; MD, mean difference; SD: standard deviation; IV: inverse variance.

Table 3

Subgroup Analyses Assessing Potential Moderating Factors for VO_2 Peak Increase in Studies on CAD Disease Included in the Meta-analysis by Population Characteristics

Research studies			Peak VO ₂				
Group	HIT groups	References	MD (95%CI)	I^2	P^{b}	P-difference ^c	
No. of participants ^a			i.	i.	i.		
< 14	4	Currie et al., ⁹ Moholdt et al., ²¹ Rognmo et al., ³² Warburton et al. ³⁴	4.12 (1.75-6.48)	0	.0006	.81	
≥ 14	5	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al. ³¹	3.77 (2.14-5.40)	42	< .00001		
BMI							
25-29.9 kg/m ²	6	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Currie et al., ⁹ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	4.46 (3.27-6.65)	0	.00001	.48	
\leq 24.9 kg/m ²	1	Kim et al. ¹⁵	6.30 (1.34-11.26)	0	< .01		
Sex							
Men	2	Rognmo et al., ³² Warburton et al. ³⁴	3.21 (-0.63-7.04)	0	.1	.71	
Men and women	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Currie et al., ⁹ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al. ²¹	3.97 (2.74-5.20)	16	< .00001		
Age, y							
> 57	4	Currie et al., ⁹ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	3.97 (2.02-5.91)	0	< .00001	.86	
≤ 5 7	4	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Wanburton et al. ³⁴	3.97 (2.02-5.91)	43	< .00001		
Methodological quality							
> 7 points	5	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Currie et al., ⁹ Madssen et al., ¹³ Moholdt et al. ³¹	3.72 (2.26-5.19)	36	< .00001	.66	
\leq 7 points	4	Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ²¹ Wanburton et al. ³⁴	4.41 (1.69-7.13)	0	.001		

95%CI, 95% confidence interval; BMI, body mass index; HIT, high-intensity interval training groups; *l*², heterogeneity; MD, mean difference; peak VO₂, peak oxygen uptake. Certain studies were not included because they did not report the value used for subgroup analysis.

^a Number of subjects of the HIT group.

^b Test for overall effect.

^c Test for subgroup differences.

Subgroup Analyses Assessing Potential Moderating Factors for VO_2 Peak Increase in Studies on HF Disease Included in the Meta-analysis by Population Characteristics

		Research studies		Pea	ak VO ₂		
Group	HIT groups	References	MD (95%CI)	I^2	P^{b}	P-difference ^c	
No. of participants ^a	î.						
< 14	6	Benda et al., ²⁴ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Koufaki et al., ³⁰ Roditis et al., ⁷ Smart et al. ³³	1.79 (0.40-3.19)	0	.01	24	
≥ 14	4	Chrysohoou et al., ⁶ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al. ¹⁷	2.87 (1.75-4.00)	72	< .00001		
Sex							
Men	2	Benda et al., ²⁴ Smart et al. ³³	1.80 (-1.01-4.61)	0		C	
Men and women	8	Chrysohoou, ⁶ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Fu et al., ¹⁶ Koufaki et al., ³⁰ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Roditis et al. ⁷	2.59 (1.68-3.50)	58	< .00001	— .b	
Age, y							
≥ 62	5	Benda et al., ²⁴ Chrysohoou et al., ⁶ Fu et al., ¹⁶ Isaksen et al., ¹⁷ Roditis et al. ⁷	2.66 (0.21-5.11)	67	.03	.73	
< 62	5	Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Huang et al., ¹⁹ Smart et al., ³³ Koufaki et al. ³⁰	2.22 (1.87-2.58)	0	< .00001	/3	
BMI							
25-29.9 kg/m ²	7	Benda et al., ²⁴ Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Isaksen et al., ¹⁷ Roditis et al., ⁷ Smart et al., ³³ Koufaki et al. ³⁰	2.04 (0.70-3.38)	14	.003	.36	
\leq 24.9 kg/m ²	3	Freyssin et al., ⁸ Fu et al., ¹⁶ Huang et al. ¹⁹	2.85 (1.72-3.99)	79	< .00001		
Methodological quality							
> 6 points	1	Chrysohoou et al. ⁶	5.00 (2.34-7.66)	_	.0002	07	
\leq 6 points	9	Benda et al., ²⁴ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Roditis et al., ⁷ Smart et al., ³³ Koufaki et al. ³⁰	2.41 (1.64-3.17)	40	< .00001	0/	

95%CI, 95% confidence interval; BMI, body mass index; HIT, high-intensity interval training groups; *l*², heterogeneity; MD, mean difference; peak VO₂, peak oxygen uptake. Certain enrolled studies were not included because the value used for subgroup analysis was not reported in them.

^a Number of subjects of the HIT group.

^b Test for overall effect.

^c Test for subgroup differences.

peak VO₂ when the duration of the protocol was < 12 weeks, with the differences being significant between the 2 subgroups (P = .05).

Regarding age, there were no statistically significant differences between subgroups (\geq 57 years and < 57 years, CAD patients [56-65 years old], and \geq 62 years and < 62 years [54-68 years], HF patients). This result indicated that age does not influence improvements in peak VO₂, although some research asserts that age is a determining factor for the occurrence of HF and CAD due to diseases associated with age.^{35,36} According to Amundsen et al.,¹² these results suggest that HIT induces changes in cardiac contractility and increases filling of the left ventricle. Thus, we found peak VO₂ improvements derived from improvements in left ventricular ejection fraction, regardless of age.

In relation to the HIT protocol, the studies suggested conducting a work ratio (work/recovery) \leq 1.33 (0.66-1.33) for patients with CAD, as it provides the same benefits as ratios > 1.33. Moreover, it would be better to use work ratios < 1 (0.5-1.33) for patients with HF. These results support some researchers' arguments that patients prefer shorter or less-intense protocols, ¹⁰ favoring adherence to the long-term protocol due to greater patient comfort.^{10,37}

Concerning work and recovery periods, there were no statistically significant differences regarding work interval duration (from 30 seconds to 4 minutes) in HF patients. Nevertheless, there were statistically significant differences in recovery intensity, as recoveries $\leq 40\%$ peak VO₂ and passive recovery did not produce adaptations in peak VO₂. The reason may be that an active recovery at greater intensity allows for the optimization of phosphocreatine resynthesis, greater lactate oxidation, and optimization of lactate neoglucogenesis.³⁸ However, although some authors have

recommended passive recoveries,¹⁰ these were not as effective as active recoveries.²⁶ For patients with CAD, there were no statistically significant differences in the work and recovery interval duration, the type of recovery, or active recovery intensity. Thus, patients with CAD had faster recoveries than HF patients. This finding could be due to CAD patients being less affected at the cardiovascular level than patients with HF, therefore having a faster recovery within the parameters mentioned.³⁹ Consequently, to improve the cardiac rehabilitation process, the design of the HIT protocol for HF patients should include active recoveries $\geq 40\%$ peak VO₂.

Regarding HIT frequency, there were no statistically significant improvements in the peak VO₂ of patients with HF when with protocols $\leq 2 \text{ d/wk} (P = .07)$,²⁴ although an improvement was found in the 3 and 4 d/wk protocols (P < .00001),^{6,7,16,17,19,27,30,33} and $\geq 5 \text{ d/wk}$ protocols (P < .01).⁸ Nevertheless, these results were not found in patients with CAD, whose peak VO₂ improved regardless of frequency (2-5 d/wk). Therefore, to obtain HIT-related peak VO₂ improvements, training frequency should be at least 3 sessions/wk for patients with HF and at least 2 sessions/wk for patients with CAD. The differences between diseases may be related to the cardiac muscle injury of HF patients, requiring a greater training frequency to obtain similar improvements compared with patients with CAD who did not have problems with the cardiac muscle.³⁷

Our systematic review and meta-analysis results indicate that peak VO₂ significantly increased (P < .00001) after HIT in patients with CAD regardless of the duration of the program. Nevertheless, there were significant differences between the protocol durations of > 12 and \leq 12 weeks (P = .05). However, for patients with HF, even though the programs > 12 weeks did not produce significant

Subgroup Analyses Assessing Potential Moderating Factors for VO₂ Peak Increase in Studies on HF Disease Included in the Meta-analysis by Exercise Characteristics

	Peak VC						
Group	HIT groups	References	MD (95%CI)	I ²	P ^a	P-difference ^b	
Number of sessions	Î.						
\geq 36 sessions	8	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Fu, ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Koufaki et al., ³⁰ Smart et al. ³³	2.69 (1.78-3.60)	56	< .00001	.26	
< 36 sessions	2	Benda et al., ²⁴ Roditis et al. ⁷	1.24 (-1.12-3.61)	0	< .00001		
Duration							
> 12 wks	2	Koufaki et al., ³⁰ Smart et al. ³³	2.45 (-0.48-5.37)	0	.1	06	
\leq 12 wks	8	Benda et al., ²⁴ Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Freyssin, ⁸ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Roditis et al. ⁷	2.53 (1.62-3.44)	59	< .00001	.90	
HIT frequency							
\geq 5 d/wk	1	Freyssin ⁸	2.90 (0.61-5.19)	-	.01	- 77	
3 or 4 d/wk	8	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Koufaki et al., ³⁰ Roditis et al., ⁷ Smart et al. ³³	2.55 (1.59-3.50)	58	< .00001	.//	
\leq 2 d/wk	1	Benda et al. ²⁴	1.30 (-2.38-4.98)	-	.49		
Session duration							
> 42 min	4	Chrysohoou et al., ⁶ Freyssin et al., ⁸ Koufaki et al., ³⁰ Smart et al. ³³	3.44 (1.95-4.94)	0	< .00001	19	
\leq 42 min	5	Benda et al., 24 Fu et al., 16 Huang et al., 19 Isaksen et al., 17 Roditis et al., 7	2.23 (1.23-3.23)	62	< .00001		
Interval work							
> 45 sec	5	Benda et al., ²⁴ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Smart et al. ³³	2.53 (1.50-3.55)	34	.002	1.00	
\leq 45 sec	5	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Koufaki et al., ³⁰ Roditis et al. ⁷	2.52 (0.89-4.15)	63	< .00001		
Interval recovery							
> 1 min	4	Benda et al., ²⁴ Fu et al., ¹⁶ Huang, ¹⁹ Isaksen et al. ¹⁷	2.46 (1.35-3.58)	73	< .00001		
$\leq 1 min$	6	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Koufaki et al., ³⁰ Roditis et al., ⁷ Smart et al. ³³	2.65 (1.27-4.03)	12	.0002	84	
HIT protocol							
HIT_L	4	Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Roditis et al. ⁷	2.45 (1.22-3.69)	7	< .00001	97	
HIT_S	6	Benda et al., ²⁴ Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Freyssin et al., ⁸ Koufaki et al., ³⁰ Smart et al. ³³	2.60 (1.37-3.82)	81	< .00001	.87	
Type of recovery							
Active (\geq 20%)	7	Benda et al., ²⁴ Freyssin et al., ⁸ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Koufaki et al., ³⁰ Roditis et al. ⁷	2.51 (1.68-3.33)	49	< .00001	.91	
Passive (< 20%)	3	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Smart et al. ³³	2.70 (-0.38-5.78)	60	.09		
Active recovery inten	sity						
$\geq 40\%$	5	Freyssin et al., 8 Fu et al., 16 Huang et al., 19 Isaksen et al., 17 Roditis et al. 7	2.60 (1.68-3.51)	58	< .00001	.59	
< 40%	2	Benda et al., ²⁴ Koufaki et al. ³⁰	1.81 (-0.88-4.50)	0	.19		
R (W/R)							
≥ 1	6	Chrysohoou et al., ⁶ Dimopoulos et al., ²⁷ Fu et al., ¹⁶ Huang et al., ¹⁹ Isaksen et al., ¹⁷ Roditis et al. ⁷	2.53 (1.45-3.62)	70	< .00001	.94	
< 1	4	Benda et al., ²⁴ Freyssin et al., ⁸ Koufaki et al., ³⁰ Smart et al. ³³	2.45 (0.83-4.07)	0	< .00001		

HF, heart failure; HIT, high-intensity interval training groups; HIT_L, protocol of HIT with long work interval; HIT_S, protocol of HIT with short work interval; l^2 , heterogeneity; MD, mean difference; peak VO₂, peak oxygen uptake; R (W/R), work-recovery ratio.

Certain studies were not included because they did not report the value used for subgroup analysis.

^a Test for overall effect.

^b Test for subgroup differences.

improvements in peak VO₂ (P = .1), there were improvements with protocols ≤ 12 weeks (P = .00001), although there were no significant differences between them (P = .96). This finding suggests that, in both diseases, the exponential improvements in peak VO₂ were produced in the first 12 weeks, when the HIT protocol was more effective. These results were probably due to the nonincremental workload included in the design of their training session, which failed to adhere to the principle of progression.⁴⁰ This is explained by the fact that when a series of effective stimuli is applied, the organism generates adaptations,

such as an increase in the muscle cross-sectional area, adaptations of energy reserves, or increased synchronization of motor units. The unchanging variability of these stimuli were therefore no longer sufficient to generate heterostasis. Thus, a coherent and progressive increase of workloads is required.⁴¹

Likewise, there were no statistical differences between patients with CAD or HF in the number of sessions (\geq 36 sessions^{6,8,11,13,16-19,21,27,30,33,34} and < 36 sessions^{7,9,15,24,31,32} [18-156 sessions for CAD patients and 18-72 sessions for HF patients]) and session duration (> 28 minutes^{11,13,15,18,21,31,32} and < 28 minutes⁹ for

Subgroup Analyses Assessing Potential Moderating Factors for VO₂ Peak Increase in Studies on CAD Disease Included in the Meta-analysis by Exercise Characteristics

			Peak VO ₂			
Group	HIT groups	References	MD (95%CI)	I^2	P ^a	P-difference ^b
Number of sessions			i			ì
\geq 36 sessions	5	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Madssen et al., ¹³ Moholdt et al., ²¹ Warburton et al. ³⁴	3.57 (1.86-5.28)	0	< .00001	.6
< 36 sessions	4	Currie et al., ⁹ Kim et al., ¹⁵ Moholdt et al., ³¹ Rognmo et al. ³²	4.25 (2.35-6.14)	35	< .00001	
Duration						
> 12 ws	3	Cardozo et al., ¹¹ Madssen et al., ¹³ Warburton et al. ³⁴	2.45 (0.62-4.29)	0	.009	.05
\leq 12 ws	6	Conraads et al., ¹⁸ Currie et al., ⁹ Kim, et al., ¹⁵ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	4.70 (3.44-5.96)	0	< .00001	
HIT frequency						
\geq 5 d/wk	1	Moholdt et al. ³¹	3.30 (0.67-5.93)	-	.01	.82
3 or 4 d/wk	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Rognmo et al., ³² Warburton et al., ³⁴ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al. ²¹	3.91 (2.48-5.33)	16	< .00001	
\leq 2 d/wk	1	Currie et al. ⁹	4.70 (1.26-8.14)	-	.007	
Session duration						
\geq 28 min	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	3.87 (2.71-5.02)	6	< .00001	.65
< 28 min	1	Currie et al. ⁹	4.70 (1.26-8.14)	-	.007	
Interval work						
\geq 4 min	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	3.90 (2.59-5.22)	16	<.00001	.99
< 4 min	2	Currie et al., ⁹ Warburton et al. ³⁴	3.92 (1.28-6.55)	0	.004	
Interval recovery						
\geq 3 min	8	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al., ³² Warburton et al. ³⁴	3.87 (2.71-5.02)	6	< .00001	.65
< 3 min	1	Currie et al. ⁹	4.70 (1.26-8.14)	_	.007	
HIT Protocol						
HIT-L	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	3.90 (2.59-5.22)	16	< .00001	.99
HIT-S	2	Currie et al., ⁹ Warburton et al. ³⁴	3.92 (1.28-6.55)	0	.004	
Type of recovery						
Active (≥20%)	8	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Rognmo et al., ³² Warburton et al., ³⁴ Kim et al., ¹⁵ Moholdt et al., ³¹ Madssen et al., ¹³ Moholdt et al. ²¹	3.87 (2.71-5.02)	6	< .00001	.65
Passive (<20%)	1	Currie et al. ⁹	4.70 (1.26-8.14)	-	.007	
Active recovery inte	nsity					
$\geq 65\%$	4	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Moholdt et al., ³¹ Moholdt et al., ²¹	4.44 (3.24-5.63)		< .00001	.27
< 65%	4	Kim et al., ¹⁵ Madssen et al., ¹³ Rognmo et al., ³² Warburton et al. ³⁴	2.86 (0.32-5.41)			
R (W/R)						
≥ 1.33	7	Cardozo et al., ¹¹ Conraads et al., ¹⁸ Kim et al., ¹⁵ Madssen et al., ¹³ Moholdt et al., ³¹ Moholdt et al., ²¹ Rognmo et al. ³²	3.87 (2.71-5.02)	6	< .00001	.65
< 1.33	2	Currie et al., ⁹ Warburton et al. ³⁴	4.70 (1.26-8.14)	_	.007	

CAD, coronary artery disease; HIT, high-intensity interval training groups; HIT_L, protocol of HIT with long work interval; HIT_S, protocol of HIT with short work interval; l^2 , heterogeneity; MD, mean difference; peak VO₂: peak oxygen uptake; R (W/R), work-recovery ratio.

Certain studies were not included because they did not report the value used for subgroup analysis.

^a Test for overall effect.

^b Test for subgroup differences.

Table 7

Recommendations on HIT Protocol for HF and CAD Patients

Disease	Frequency, d-wk	Duration program, wks	Session duration, min	Intensity of recovery, peak VO ₂ , $\%$	Ratio, work/recovery
HF	≥ 3 (2-5)	≥ 6 (6-24)	30-60 (28-60)	$\geq 40\%$ (40-70%)	\geq 1.33 (0.66-1.33)
CAD	≥ 2 (2-5)	≥ 6 (4-52)	30-60 (28-60)	$\geq 40\%$ (0-70%)	≥ 1 (0.5-1.33)

CAD, coronary artery disease; HF, heart failure; HIT, high-intensity interval training groups; peak VO₂: peak oxygen uptake.

and $HF^{6,16,19}$ patients. These results, together with those obtained in this meta-analysis, suggest that the maximum benefits in peak VO_2 would be obtained between weeks 6 and 12. There is therefore a need to accomplish an adequate training period to perceive the effects produced in peak VO_2 by long-term training protocols. For this reason, it is important to optimize training through adherence to the general principles of training.

In relation to the HIT protocol, this meta-analysis confirms the results that both HIT protocols with short work intervals and protocol of HIT with long work intervals produce significant improvements in peak VO_2 in patients with $CAD^{9,14,18}$ and in patients with HF.^{6,8,16,19} Likewise, no significant differences were found between performing an HIT protocol with short work interval or long work interval in either disease. These results may be due to the improvements in peak VO₂ of cardiovascular diseases being more closely related to the intensity rather than to the duration of the HIT protocol.⁵ This is because intensity is the parameter that produces greater adaptations in the cardiovascular system such as an increase in muscle cross-sectional area, adaptations of energy reserves, or increased synchronization of motor units.^{33,34} Therefore, the type of protocol is not as important in the peak VO₂ improvement as the training intensity and frequency, for CAD and HF patients.

According to our meta-analysis, HIT improved peak VO₂ in patients with CAD or HF, although there were statistically significant differences between these diseases with a greater improvement in patients with HF (P = .03). These results may be due to the initial differences in peak VO₂ (16.2-31.8 mL/kg/min⁻¹ in CAD patients and 10.7-19.1 mL/kg/min⁻¹ in HF patients), derived from the fact that HF patients have impaired cardiac function and the cardiac muscle was not able to pump enough blood to supply the tissues.³² However, CAD patients saw a lesser impact on their peak VO₂, as they probably had better myocardial contractility, particularly if they had undergone coronary intervention.⁴²

As has been previously suggested, this should allow the possibility of a higher increase in cardiac output because of a higher systolic volume, resulting in a greater increase in left ventricular ejection fraction in HF patients due to their having a lower threshold of adaptation to the stimulus.⁷ Similarly, this improvement could be caused by a larger relative increase in exercise-induced vasodilation, hemoglobin, the skeletal muscle oxidative capacity,^{5,8,18} or by the sum of all these factors.

Although it was not an objective of this study, there are studies that compare the effects on peak VO₂ between HIT and continuous training. Although most studies have suggested that HIT is superior to continuous exercise, ^{15,16,21,23,43} some studies have reported that HIT was not superior to continuous exercise programs in relation to peak VO₂ in the 2 diseases.^{5,7,9,17–20,44,45}

The present study makes an important contribution to the understanding of the effectiveness of HIT training programs in heart diseases. Thus, this study provides evidence for the potential applicability of HIT training programs as part of the treatments used for CAD and HF.

Practical Application

The results of this study indicate that HIT positively affects peak VO_2 in people with HF or CAD, providing greater advantages in HF

patients. These findings could be used by physicians, physical conditioning trainers and heart rehabilitation teams to develop specific training programs in order to optimize the functioning of the patient's heart. However, other variables have to be taken into account, such as age, training frequency, the duration of program, and the type of recovery. The HIT program should therefore be adapted to the individual characteristics of each patient. The recommended dosage for each disease is shown in Table 7.

Limitations

The main limitations of this study are as follow: a) the randomized controlled trials did not use the same methods to control the intensity of the training sessions; b) the protocols and the age of the participants were widely heterogeneous; c) most of the included studies had a sample with few patients; and d) in HIT protocols, one aspect is the design and another the intensity that patients are capable of achieving. Therefore, the possibility that a patient has not been able to accomplish the planned intensities should be considered.

CONCLUSIONS

The results of this study show that HIT is an effective method for the treatment of HF and CAD by improving peak VO₂, with the increase being significantly higher in HF patients. To optimize these benefits, recovery intervals should be active at intensities between 40% to 60% of peak VO₂ for HF patients, and the frequency should be ≥ 2 d/wk for CAD patients and ≥ 3 d/wk for HF patients. This study opens a new line of research that could be used to optimize high-quality exercise training protocols in an effort to develop the most effective and efficient method for the treatment of heart disease and other diseases.

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CONFLICTS OF INTEREST

None declared.

WHAT IS KNOWN ABOUT THE TOPIC?

- The HIT produces positive cardiovascular adaptations in HF and CAD patients.
- The HIT improves peak VO₂ in HF and CAD patients.

WHAT DOES THIS STUDY ADD?

- The HIT is more effective in improving peak VO₂ in HF patients than in CAD patients.
- To obtain improvements in peak VO₂ through HIT, \geq 3 sessions per week for at least 6 weeks are needed for HF patients, and \geq 2 sessions per week for at least 6 weeks for CAD patients.
- Active recovery at intensities between 40% to 60% of peak
 VO₂ should be used to improve peak VO₂ in HF patients.

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