Optimal aerobic exercise intensity and its influence on the effectiveness of exercise therapy in patients with pulmonary arterial hypertension: a systematic review

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Background: Exercise intensity in exercise training programs is an important determinant of program efficacy, such as improvement in exercise capacity and quality of life (QOL). It is not well known whether differently applied exercise intensities are efficacious when used in exercise-based cardiac rehabilitation programs for patients with pulmonary arterial hypertension (PAH).

Methods: Three databases (PubMed, EMBASE, and CINAHL) were searched with the following inclusion criteria: comparative study of exercise interventions for patients with pulmonary arterial hypertension. Three clinical specialists (a physician, nurse, and exercise physiologist) selected the included articles using the process of systematic review. Included articles were grouped according to aerobic exercise intensity: low, moderate-to-vigorous, and vigorous. The level of evidence for each study was rated using Sackett’s levels of evidence.

Results: Of 1,452 studies reviewed, 8 were included according to the inclusion criteria (3 randomized controlled trials (RCTs), 3 prospective studies, and 2 case series). Exercise capacity for a six-minute walk distance (mean: 57.7 m) and QOL improved in the above moderate intensity group, while the low intensity group did not show improvement after intervention. For termination criteria, data obtained from the reviewed articles were not sufficient to suggest any exercise intensity recommendations for patients with pulmonary arterial hypertension.

Discussion: The findings in this study suggest that at least moderate aerobic exercise intensity is needed to significantly improve six-minute walk distance and QOL in individuals diagnosed with World Health Organization Group 1 of pulmonary arterial hypertension. There is a need for prospective RCTs comparing different exercise intensities in this patient population.

Keywords: Pulmonary arterial hypertension (PAH); exercise tolerance; quality of life (QOL); exercise therapy

doi: 10.21037/jtd-20-3296
View this article at: https://dx.doi.org/10.21037/jtd-20-3296
Introduction

Pulmonary arterial hypertension (PAH) is a rare and progressive disease characterized by the progressive development of dyspnea and fatigue with increasing deterioration of exercise tolerance (1,2). Treatment with medication is a first option to reduce mortality and to improve dyspnea (3). Exercise therapy has been suggested as a treatment option for patients with PAH since 2000, and the clinical evidence for this has been increasing (4).

An evidence-based randomized controlled study on exercise effects in patients with pulmonary hypertension was reported for the first time in 2006 (5), and a meta-analysis that compared the effects of exercise therapy was published in 2015 (6). Several studies (5,7-9) demonstrated that exercise therapy for PAH patients contributed to improved exercise capacity, quality of life (QOL), muscle strength, and respiratory muscle strength. The exercise types typically included aerobic exercise, resistance training, and respiratory muscle strength exercise (10).

Considering the safety and efficacy of exercise therapy for patients with PAH, optimal exercise intensity is an important factor to prevent adverse events and to increase the effectiveness of exercise therapy (11-13). High intensity exercise can cause severe adverse events including arrhythmias, syncope, and respiratory disease in 13% of these patients (11).

Exercise intensity was usually prescribed and monitored using heart rate (HR), pulse oxymetry saturation (SpO₂), and rating of perceived exertion (RPE) in PAH patients (12,14-17). Some criteria including HR below 120 bpm and SpO₂ below 85% were used as termination criteria during exercise training in individuals with PAH (15,17-21).

However, the majority of review articles (6,7,10,13) have focused their investigations on the effectiveness of exercise interventions with respect to exercise capacity, muscle strength, and QOL. The improvement of exercise tolerance may be different according to the applied exercise intensity. Applying different exercise intensities is associated with different levels of effectiveness in endothelium-dependent vasodilation, depending on the interaction between the intensities of both exercises (22).

Therefore, the objective of this review was to investigate the effects of exercise-based cardiac rehabilitation programs according to exercise intensity to improve exercise capacity and QOL, and to describe the termination criteria used during exercise training in patients with World Health Organization (WHO) Group 1 of pulmonary arterial hypertension.

We present the following article in accordance with the PRISMA reporting checklist (available at https://dx.doi.org/10.21037/jtd-20-3296).

Methods

Protocol and registration

The protocol was registered with PROSPERO (PROSPERO2020: CRD42020184937). The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines were followed.

Eligibility criteria

The following PICOS (Participants, Interventions, Comparison, Outcome, and Study Design) criteria were used to generate our research:

(I) Participants: patients with WHO Group 1 of PAH;
(II) Intervention: exercise training;
(III) Comparison: exercise intensity, termination criteria;
(IV) Outcome: exercise capacity, QOL;
(V) Study design: randomized controlled trials (RCTs), prospective studies, and observational studies.

Search strategy

This literature review search was performed using the PubMed, EMBASE, and CINAHL databases, which were searched using the following terms: (“Exercise”[Mesh] or “Exercise Therapy”[Mesh] or “Cardiac Rehabilitation”[Mesh] or “Resistance Training”[Mesh]) and (“Hypertension, Pulmonary”[Mesh] or “Pulmonary Arterial Hypertension (PAH)”[Mesh]). The search included articles published between 1980 and June 2020, and all articles identified in the search were evaluated.

Criteria for inclusion

Studies were included if they met the following criteria: RCTs or prospective observational studies or case series with comparisons between before and after exercise therapy. The participants were diagnosed as WHO Group 1, defined as PAH. The studies with case reports and literature reviews and trials that included other WHO group of pulmonary hypertension groups were excluded. Articles written in languages other than English were also excluded.
Study selection

Two reviewers (YGS and SKO) independently reviewed the studies and extracted articles that met the inclusion criteria according to the review protocol. They screened the articles by checking titles and abstracts first, and then full texts. A third reviewer (JS) reviewed the full text of an article to determine whether it should be included in cases where the original two reviewers had disagreements about those articles.

Quality of selected articles

The quality of each study was assessed using Sackett’s level of evidence scale from the strongest (rating =1) to weakest (rating =5), where ranked RCTs are considered the highest level and case series or expert opinions are considered the lowest level (Table 1).

Statistical analysis

Due to the heterogeneity and non-uniformity of the data in the included studies, the results are summarized in a descriptive manner.

Results

Study selection

In total, 1,452 articles were found according to the process of this study, and the results are as follows. The numbers of article types (RCTs, prospective studies, case series) found in PubMed, EMBASE, and CINAHL were 637, 684, and 131, respectively. Eight articles from the 1,452 were selected. Three were RCTs, 3 were prospective studies, and 2 were case series (Figure 1).

Study characteristics

The most common types of PAH were idiopathic and connective tissue disease. Most of the patients had functional classifications from II to III, and only three trials (9,14,16) included patients with functional classification IV. A summary of the included studies has been provided in Table 2. The common inclusion criteria were adults aged 18 and older, medically stable condition, and no change in medication therapy for 3 months before participation.

Classification of aerobic exercise intensity

Heart rate domains were used in the majority of studies to prescribe target exercise intensity, and %peak oxygen uptake (peak VO₂) was used in one study (19). Aerobic exercise intensity ranged from 50% to 80% of %heart rate reserve (HRR) or %maximal heart rate (HRmax) or %peak VO₂. For the purposes of this review article, exercise intensity was categorized into 3 different groups according to American College of Sports Medicine (ACSM) guidelines (Table 3) (23); the low-intensity exercise group required less than 120 bpm or a rating perceived exertion (RPE) of 13 to perform (18,20), the moderate to vigorous-intensity exercise group required 60–80% of HRmax or 50–70% of peak VO₂ (9,12,19,21), and the vigorous-intensity exercise group required 70–80% of HRR (14,16) to perform.

Table 1 Evidence level according to Sackett’s levels of evidence

<table>
<thead>
<tr>
<th>Level</th>
<th>Explanation of each level</th>
<th>Reviewed article’s level</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A</td>
<td>Systematic Review of Randomized Controlled Trials (RCTs)</td>
<td>–</td>
</tr>
<tr>
<td>1B</td>
<td>RCTs with Narrow Confidence Intervals</td>
<td>3</td>
</tr>
<tr>
<td>1C</td>
<td>All or None Case Series</td>
<td>–</td>
</tr>
<tr>
<td>2A</td>
<td>Systematic Review Cohort Studies</td>
<td>–</td>
</tr>
<tr>
<td>2B</td>
<td>Cohort Study/Low Quality RCT</td>
<td>–</td>
</tr>
<tr>
<td>2C</td>
<td>Outcomes Research</td>
<td>3</td>
</tr>
<tr>
<td>3A</td>
<td>Systematic Review of Case-Controlled Studies</td>
<td>–</td>
</tr>
<tr>
<td>3B</td>
<td>Case-controlled Study</td>
<td>–</td>
</tr>
<tr>
<td>4</td>
<td>Case Series, Poor Cohort Case-Controlled</td>
<td>2</td>
</tr>
<tr>
<td>5</td>
<td>Expert Opinion</td>
<td>–</td>
</tr>
</tbody>
</table>
Exercise capacity

The six-minute walk test (6MWD) and cardiopulmonary exercise test (CPET) are conducted to evaluate exercise capacity. All studies reported the results of the 6MWD, while the CPET was not conducted in two trials (9,12). The baseline 6MWD was reported to range from 375±92 to 496±108 m in the reviewed articles. The improvement seen in the 6MWD ranged from 37 to 81 m, and the median value was 56 m. In the low intensity group, one study did show non-significant change (18) and others reported an improvement of 39 m (20). The moderate to vigorous group reported improvements in the 6MWD of 37 m (19), 71 m (21), 81 m (9), and 67 m (12). Changes in the vigorous-intensity group were 56 m (14) and 53 m (16). The CPET was conducted with a treadmill in three studies and with a cycling ergometer in another three studies. At baseline, maximal oxygen uptake ranged from the lowest level of 11.4±2.2 mL/kg/min (12) to the highest level of 17.6 mL/kg/min (14). The improvement in peak VO$_2$ showed a range from 0.3 to 2.3 mL/kg/min, and the highest improvement of 14% was reported in the study conducted by Grünig et al. (21). Two trials (9,20) did not conduct the CPET, and one trial did not report the results of post-intervention evaluations (Table 4).

QOL

The 36-Item Short Form Health Survey (SF-36) and the Cambridge Pulmonary Hypertension Outcome Review (CAMPION) questionnaire were used to evaluate QOL in the reviewed articles. Of two studies in the low-intensity group, one did not evaluate QOL and the other showed no significant difference in the SF-36 domains. In the moderate to vigorous-intensity group, three studies (12,19,21) reported the results of QOL, but one trial (9) did not measure QOL. Of these articles, one article (12) showed a significant improvement in the bodily pain domain (P=0.05). A study reported by Grünig et al. (21) showed that the SF-36 domains including physical function, general health, vitality, social function, and mental health demonstrated significant improvements. Another study (10) did not report a significant difference after an intervention of 10 weeks. In the vigorous-intensity group, one trial (14) of two articles reported a significant improvement in SF-36 domains including physical function, role physical, general health, vitality, social function, and mental health and the other study did not conduct an evaluation of QOL (16). The CAMPION questionnaire was only used in a study conducted by Chan et al. (14), who reported an improvement in the domains of QOL, symptoms, and energy. Mental domains showed greater improvement when compared to the physical domains for the SF-36 questionnaire (12,14,21).

Criteria for terminating exercise

For safety in exercise training, four of the eight reviewed
Table 2 A summary of the characteristics of included studies and exercise prescriptions

<table>
<thead>
<tr>
<th>Author (year)/design</th>
<th>Subjects</th>
<th>Types of PAH</th>
<th>Functional classification</th>
<th>Aerobic exercise</th>
<th>Resistance training</th>
<th>Respiratory training</th>
</tr>
</thead>
<tbody>
<tr>
<td>de Man et al. [2009] (18)/Prospective study</td>
<td>N=19 (male: 4, female: 15); Age: 42.0±13.0</td>
<td>Idiopathic: 19</td>
<td>NYHA: Class II, 3; Class III, 16</td>
<td>F: 3/week; T: Cycling; T: 20–25 min; D: 12 weeks</td>
<td>F: 3/week; I: 50–75% of 1RM for strength, 30–40% for endurance; T: NR; T: NR; D: 12 weeks</td>
<td>None conducted</td>
</tr>
<tr>
<td>Butane et al. [2019] (20)/Case series</td>
<td>N=6 (male: 1, female: 5); Age: 68.0±7.6</td>
<td>Idiopathic: 6</td>
<td>NYHA: I, 1; II, 3; III, 2</td>
<td>F: 3/week; T: walking; T: 20–40 min; D: 12 weeks</td>
<td>F: 5/week; I: 5–10 rep; T: to 6–7 resistance; T: 20–40 min; D: 12 weeks</td>
<td>None conducted</td>
</tr>
<tr>
<td>Kabitz et al. [2014] (9)/Case series</td>
<td>N=7 (male: 3, female: 4); Age: 59.6±11.1</td>
<td>Idiopathic: 5; CTD: 2</td>
<td>WHO: Class III, 6; Class IV, 1</td>
<td>F: 7/week; T: cycling; T: 10–25 min; D: 15 weeks</td>
<td>F: 5/week; I: 500 to 1,000 g; T: dumbbell training; T: 30 min; D: 15 weeks</td>
<td>None conducted</td>
</tr>
<tr>
<td>Becker-Grünig et al. [2013] (12)/Prospective study</td>
<td>N=20 (male: 4, female: 16); Age: 48.0±11.0</td>
<td>ACHD</td>
<td>WHO: Class II, 6; Class III, 14</td>
<td>F: 7/week; T: cycling; T: 30–45 min; D: 15 weeks</td>
<td>F: 5/week; I: 500 to 1,000 g; T: dumbbell training; T: 30 min; D: 15 weeks</td>
<td>None conducted</td>
</tr>
<tr>
<td>Karapolat et al. [2019] (19)/RCT</td>
<td>N=24 (EG: 12, CG: 12), (male: 11, female: 13); Age (EG: 34, CG: 40)</td>
<td>Idiopathic: 3; Congenital: 11; Rheumatologic disease: 4</td>
<td>WHO: Class II, 18; Class III, 6</td>
<td>F: 3/week; T: treadmill walking; T: 30 min; D: 8 weeks</td>
<td>None conducted</td>
<td>F: 3/week; I: NR; T: breathing exercise; T: NR; D: 8 weeks</td>
</tr>
<tr>
<td>Grünig et al. [2012] (21)/Prospective study</td>
<td>N=21 (male: 1, female: 20); Age: 52.0±18.0</td>
<td>Systemic sclerosis: 8; SLE: 7; MCTD: 2; Other: 4</td>
<td>WHO: II, 9; III, 7; IV, 5</td>
<td>F: 7/week; T: bicycle ergometer; T: 30–45 min; D: 15 weeks</td>
<td>F: 5/week; I: 500 to 1,000 g; T: dumbbell training; T: 30 min; D: 15 weeks</td>
<td>None conducted</td>
</tr>
<tr>
<td>Chan et al. [2013] (14)/RCT</td>
<td>N=23 (EG: 10, CG: 13), female: 23; Age (EG: 53.0±13.0, CG: 55.5±8.5)</td>
<td>Idiopathic: 5; Drug-induced: 1; CTD: 17</td>
<td>WHO: Class I, 1; Class II, 12; Class III, 9; Class IV, 1</td>
<td>F: 24–30 sessions; T: treadmill walking; T: 30–45 min; D: 10 weeks</td>
<td>None conducted</td>
<td>None conducted</td>
</tr>
<tr>
<td>Weinstein et al. [2013] (16)/RCT</td>
<td>N=24 (EG: 11, CG: 13), female: 24; Age (EG: 53.4±12.4, CG: 55.3±8.7)</td>
<td>Idiopathic: 6; MCTD: 1; RA: 1; Scleroderma: 12; Sjogren's syndrome: 2; SLE: 2</td>
<td>WHO: Class I, 2; Class II, 16; Class III, 15; Class IV, 2</td>
<td>F: 3/week; T: treadmill walking; T: 30–45 min; D: 10 weeks</td>
<td>None conducted</td>
<td>None conducted</td>
</tr>
</tbody>
</table>

ACHD, adult congenital heart disease; WHO, World Health Organization; CPET, cardiopulmonary exercise test; MHR, maximal heart rate; P, protocol; M, mobility; F, frequency, I, intensity, T, type; T, time; D, duration; RCT, randomized controlled trial; EG, exercise group; CG, control group; CTD, connective tissue disease; NYHA, New York Heart Association; HRR, heart rate reserve; MWT, minute walk test; RPE, rating of perceive exertion; R, repetition; RM, repetition maximum; RA, rheumatic arthritis; MCTD, mixed connective tissue disease; SLE, systemic lupus erythematosus; TMT, treadmill test; VO₂R, oxygen uptake reserve.
articles suggested the criteria for terminating exercise as heart rates more than 120 bpm (18,20) or 130 bpm (9), less than 85% (9,18,20) of $\text{SpO}_2$, or >6 on the Borg scale (20).

**Discussion**

This study showed that moderate or higher intensity aerobic exercise was associated with an improvement in exercise capacity and QOL and that HR and $\text{SpO}_2$ were most commonly used for the termination criteria.

**Classification of aerobic exercise intensity**

A proper definition of aerobic exercise intensity is important to enhance the effectiveness of exercise therapy and to prevent the occurrence of adverse events during exercise (24). Cardiovascular adaptations to training are intensity-dependent, and exercise intensity has emerged as an important variable in clinical investigations (22,25). Exercise intensity domains for aerobic exercise include $\%\text{HR}_{\text{max}}$, $\%\text{HRR}$, $\%\text{peak VO}_2$, and $\%\text{VO}_2\text{max}$ (26). Some systematic review articles reported that recommended aerobic exercise intensity is 60 to 80 $\%\text{peak VO}_2$ or $\%\text{HRR}$ (7,13). In this study, the common domains used for exercise intensity were $\%\text{peak VO}_2$, $\%\text{HR}_{\text{max}}$, and $\%\text{HRR}$. According to ACSM guidelines (23), exercise intensity is classified into low-intensity physical activity at 20–39%, moderate-intensity activity at 40–59%, and high-intensity activity at 60–84% of peak VO₂ or HRR. When expressed as a percentage of the HRmax, low-intensity activity is 40–63%, moderate-intensity activity is 64–76%, and vigorous-intensity activity is 76–90%. We classified these reviewed articles into 3 different groups based on ACSM guidelines: a low-intensity group, a moderate to vigorous-intensity group, and a vigorous-intensity group. In the low-intensity group, the exercise capacity and QOL did not show significant improvement after the exercise intervention. The results in the moderate to vigorous-intensity group were similar to those in the vigorous-intensity group. This finding differs from previous studies (5,27) that recommended low-intensity exercise might be a beneficial intervention to improve clinical outcomes in PAH patients. Therefore, further study with high quality RCTs is required to confirm the effects of different exercise intensities on exercise tolerance and QOL in patients diagnosed with PAH.

**Exercise capacity**

The six-minute walk test in PAH patients has been widely used to measure exercise capacity and to predict prognoses (28). In the reviewed articles, the 6MWD was reported in all trials, and the highest improvement was shown to be 81 m in the moderate to vigorous-group. The range of improvement in the 6MWD was 31 to 81 m; this is similar to the result of a systematic review, which reported an improvement range from 17 to 96 m in patients with PAH (7). The mean improvement of 6MWD was 57.7 m in this study, a greater improvement than that reported in a previous study, which was a mean improvement of 33.8 m for those in WHO Group 1 (29). Another previous study reported that an increase in 6MWD of over 41 m may be the minimally important distance for clinical significance (30). On the basis of this report, two studies in the low-intensity group and one in the moderate to vigorous-intensity group did not achieve the improvement of a clinically significant distance of ≥41 m. Two RCT studies in the vigorous-intensity group showed increases in the 6MWD of over 41 m. The differences among the groups might be explained as follows:

<table>
<thead>
<tr>
<th>Exercise intensity</th>
<th>$\text{VO}_2\text{R (%)}$; $\text{HRR (%)}$</th>
<th>Maximal HR (%)</th>
<th>12 METs; $\text{VO}_2\text{max}$</th>
<th>10 METs; $\text{VO}_2\text{max}$</th>
<th>8 METs; $\text{VO}_2\text{max}$</th>
<th>6 METs; $\text{VO}_2\text{max}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Very light</td>
<td>&lt;20</td>
<td>&lt;50</td>
<td>&lt;3.2</td>
<td>&lt;2.8</td>
<td>&lt;2.4</td>
<td>&lt;2.0</td>
</tr>
<tr>
<td>Light</td>
<td>20–39</td>
<td>50–63</td>
<td>3.2–5.3</td>
<td>2.8–4.5</td>
<td>2.4–3.7</td>
<td>2.0–3.0</td>
</tr>
<tr>
<td>Moderate</td>
<td>40–59</td>
<td>64–76</td>
<td>5.4–7.5</td>
<td>4.6–6.3</td>
<td>3.8–5.1</td>
<td>3.1–4.0</td>
</tr>
<tr>
<td>Hard</td>
<td>60–84</td>
<td>77–93</td>
<td>7.6–10.2</td>
<td>6.4–8.6</td>
<td>5.2–6.9</td>
<td>4.1–5.2</td>
</tr>
<tr>
<td>Very hard</td>
<td>&gt;85</td>
<td>&gt;94</td>
<td>&gt;10.3</td>
<td>&gt;8.7</td>
<td>&gt;7.0</td>
<td>&gt;5.3</td>
</tr>
<tr>
<td>Maximal</td>
<td>100</td>
<td>100</td>
<td>12</td>
<td>10</td>
<td>8</td>
<td>6</td>
</tr>
</tbody>
</table>

$\text{VO}_2\text{R}$, oxygen uptake reserve; $\text{HRR}$, heart rate reserve; HR, heart rate; METs, metabolic equivalents; $\text{VO}_2\text{max}$, maximal oxygen uptake.

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**Table 3 Classification of physical activity intensity recommended by ACSM’s guidelines**
<table>
<thead>
<tr>
<th>Author (year)/ design</th>
<th>Groups</th>
<th>Exercise intensity</th>
<th>6MWD (m)</th>
<th>Peak VO$_2$ (ml/kg/min)</th>
<th>Quality of life (score)</th>
</tr>
</thead>
<tbody>
<tr>
<td>de Man et al. [2009] (18)/ Prospective study</td>
<td>Low-intensity group</td>
<td>Less than 120 bpm or RPE 13</td>
<td>Pre: 496±108; Post: non-significant; Exercise time: 8%; (P&lt;0.001)</td>
<td>M: Cycling; Peak VO$_2$; Pre: 15±4; Post: none reported</td>
<td>None conducted</td>
</tr>
<tr>
<td>Butane et al. [2019] (20)/Case series</td>
<td>Low-intensity group</td>
<td>Less than 120 bpm or RPE 13</td>
<td>Pre: 375±92; Post: 408±94; 39±17.5 (P=0.04). Improved 8%</td>
<td>None conducted</td>
<td>Type: SF-36. No significant difference</td>
</tr>
<tr>
<td>Kabitz et al. [2014] (9)/Case series</td>
<td>Moderate to vigorous-intensity group</td>
<td>60–80% of HRmax</td>
<td>Pre: 417±51; 3 weeks: 509±39; Post: 498±39; 81±30 (P&lt;0.001). Improved 16%</td>
<td>None conducted</td>
<td>None conducted</td>
</tr>
<tr>
<td>Becker-Grüning et al. [2013] (12)/ Prospective study</td>
<td>Moderate to vigorous-intensity group</td>
<td>60–80% of HRmax</td>
<td>Pre: 423±80; 3 weeks: 486±93; Post: 466±102; 67±59 (P=0.001). Improved 16%</td>
<td>M: Cycling; Pre: 11.4±2.2; 3 weeks: 12.4±2.2; Post: 12.3±2.4; Improved 11%</td>
<td>Type: SF-36. Only BP was significantly different (62.2±33 to 82.4±20; P=0.05)</td>
</tr>
<tr>
<td>Karapolat et al. [2019] (19)/RCT</td>
<td>Moderate to vigorous-intensity group</td>
<td>50–70% of peak VO$_2$</td>
<td>Pre: 390; Post: 427; 37 (P=0.32). Improved 9%</td>
<td>M: Treadmill; Pre: 15.1; Post: 15.4; (P=0.06). Improved 2%</td>
<td>Type: SF-36. No significant difference</td>
</tr>
<tr>
<td>Grüning et al. [2012] (21)/ Prospective study</td>
<td>Moderate to vigorous-intensity group</td>
<td>60–80% of HRmax</td>
<td>Pre: 386±121; 3 weeks: 425±118; Post: 447±139; 71 (P=0.03). Improved 14%</td>
<td>M: Cycling; Pre: 11.8±3.4; 3 weeks: 13.6±3.4; Post: 14.1±3.5; (P=0.008). Improved 14%</td>
<td>Type: SF-36; PF 33.5±20 to 45.2±20 (P=0.029); GH 31.6±15 to 35.9±14 (P=0.049); Vitality 38.1±14 to 48.1±18 (P=0.021); SF 61.8±13 to 71.4±23 (P=0.008); MH 56.8±21 to 66.2±16 (P=0.033)</td>
</tr>
<tr>
<td>Chan et al. [2013] (14)/RCT</td>
<td>Vigorous-intensity group</td>
<td>70–80% of HRR</td>
<td>Pre: 411±73; Post: 467±86; 56 (P=0.002). Improved 12%</td>
<td>M: Treadmill; Pre: 17.6; Post: 18.9; Improved 7%</td>
<td>Type: SF-36; PF (P=0.009), RP (P=0.023), GH (P=0.002), Vitality (P=0.016), MH (P=0.028). Type: CAMPHOR: QOL (P=0.003), symptoms (P=0.005), energy (P=0.008), breathlessness (P=0.041), mood (P=0.032)</td>
</tr>
<tr>
<td>Weinstein et al. [2013] (16)/RCT</td>
<td>Vigorous-intensity group</td>
<td>70–80% of HRR</td>
<td>Pre: 412±69; Post: 465±113; 53±44 (P=0.003). Improved 12%</td>
<td>M: Treadmill; Pre: 97.5±41 W; Post: 99.6±42 W; 25±22 W (P=0.003). Improved 2%</td>
<td>None conducted</td>
</tr>
</tbody>
</table>

6MWD, six-minute walk distance; RPE, rating perceived exertion; M, mode; SF, short-form; HRmax, maximal heart rate; HRR, heart rate reserve; W, watt; RCT, randomized controlled trial; PF, physical function; GH, general health; BP, bodily pain; SF, social function; MH, mental health; RP, role physical; CAMPHOR, Cambridge Pulmonary Hypertension Outcome Review; QOL, quality of life.
by the different exercise durations and frequencies applied in each study, which may have affected the results. An RCT applying the same exercise duration and frequency is required to confirm differences between groups according to different exercise intensities. Another clinical outcome regarding exercise capacity is the maximal oxygen uptake measured by CPET, and the improvement, in the present study, is similar to the results (1.1 to 2.1 mL/kg/min) reported by Babu et al. (7). Of the reviewed articles, we were not able to compare the results between three trials, because two trials did not conduct the CPET and one did not report post intervention results (18). In the low-intensity group, maximal oxygen uptake was not reported. The highest change (2.3 mL/kg/min) in peak VO₂ was revealed in the moderate to vigorous-intensity group with 60–80% of HRmax, and the mean improvement was 7% in three studies. The vigorous-intensity group showed a mean increase of 4.5%. The mean value was higher in the moderate to vigorous-intensity group than in the other intensity groups and the reason for this needs to be investigated in future studies.

QOL

Patients with PAH have anxiety and depression, and this could lead to a decrease in QOL (31). The generic SF-36 questionnaire was used in most of the studies, and CAMPHOR was used for measuring disease-specific conditions (32). A decreased QOL is associated with an increased mortality in patients with PAH (33). In the reviewed articles, the generic SF-36 was commonly used to measure QOL and only one study used a disease-specific CAMPHOR questionnaire. In particular, PAH patients report a reduced QOL in the physical components of the SF-36 domains (34,35). This review study showed that the physical components score was lower than the mental health components score.

One study in the low-intensity group (20) did not show a significant change after an exercise intervention of 12 weeks. Additionally, a study (19) conducted for 8 weeks with a moderate to vigorous-intensity of 50–70% of peak VO₂ showed no significant difference in QOL pre- post-training. The exercise frequency in the two studies was 3 times per week, and this frequency was lower than other studies (14,21) that demonstrated a significant difference in QOL pre- post-training. Three of the eight studies showed a statistically significant difference in SF-36 domains including bodily pain, physical function, general health, vitality, social function, and mental health pre- post-training. The studies had different levels of improvement in 6MWD, and the study conducted by Grünig et al. (21) reported an increase of 71 m after the exercise intervention. Another study (21) also showed the highest significant improvement in the SF-36 domains, while one study with an RCT with the lowest improvement in 6MWD (37 m) showed no significant difference in QOL scores pre- post-training. The difference among the moderate to vigorous intensity-groups may be explained by the correlation between the 6MWD and QOL. That is, improvement in the 6MWD is associated with increased QOL in individuals with PAH (36,37). Study design also should be considered as a reason for this difference, because the prospective study and RCT showed different results despite applying the same exercise intensity. For the vigorous-intensity groups included in two studies, one study reported a significant change in the SF-36 domains including physical function, role physical, general health, vitality, social function, and mental health. Two studies were showed similar results for the 6MWD, but two results could not be compared among the studies, because one study did not evaluate QOL.

The CAMPHOR questionnaire is a disease-related questionnaire for patients with PAH and has superior psychometric properties, compared with the SF-36 (38). The specific disease-related CAMPHOR was only reported in a study conducted by Chan et al. (14) of our reviewed articles. Therefore, further study is needed to measure QOL using the CAMPHOR questionnaire in PAH patients to confirm the relationship between the disease-specific questionnaire and QOL.

Termination criteria

Timing for exercise termination is an important factor to consider to prevent adverse events due to participation in an exercise program (11,12). Some adverse events that included respiratory and gastrointestinal infections were reported in the reviewed articles (12,21). Several parameters such as HR, SpO₂, and RPE were used to monitor patient state in four studies (9,15,18,20). The criterion for an HR upper limit of 120 bpm has commonly been used as an indication for termination, and it was also recommended as a target HR for inpatient cardiac patients (22). Hypoxemia may contribute to a sensation of dyspnea by predisposing the respiratory muscles to fatigue (39). Portable pulse oximetry is commonly used to measure oxygen saturation during exercise. The termination criterion with respect
to \( \text{SpO}_2 \) was defined as when the \( \text{SpO}_2 \) dropped to less than 85% or 88% in several studies (9,15,18,20). Oxygen saturation \( \leq 88\% \) is referred to as resting hypoxemia, and maintaining this criterion is thought to be an important part of exercise in PAH patients (40). Another termination criterion was Borg’s scale (10-point scale) for RPE, and dyspnea ratings between 3 and 5 have been recommended as the proper exercise intensity in patients with COPD (40). In this review study, only one study by Butâne et al. (20) used Borg’s scale to monitor the severity of dyspnea and recommended a rating of more than 6 as a termination criterion. However, the majority of the reviewed studies did not suggest termination criteria, and there was no consensus among the studies on what such criteria should consist of. Therefore, more strict termination criteria are needed to prevent adverse events or for safety during exercise training in patients from WHO Group 1.

**Limitations**

There are several limitations in this review article. First, only 3 large databases, PubMed, EMBASE, and CINAHL, were searched, despite the existence of some other large and relevant databases such as CENTRAL and PEDro. Second, the subjects enrolled in the present study were patients in WHO Group 1. In addition, there were lack studies on low intensity exercise in reviewed articles since only WHO group 1 in pulmonary hypertension is included. Therefore, it is not suitable to apply the results of this trial to other PH groups. A future study is needed to confirm differences in the effects of different exercise intensities in each PH group. Third, the reviewed articles did not include sufficient RCTs. That is, the evidence level is poor with respect to confirming the effectiveness of different exercise intensities in individuals with PAH. Large clinical trials designed as RCTs with long-term follow up are required to determine the role of exercise intensity in exercise programs for PAH patients.

**Conclusions**

The main findings in this study suggest that aerobic exercise at moderate or higher intensity is needed to improve exercise tolerance and QOL in PAH patients, but the quality of evidence is low. Additional RCTs are needed to compare the effects of different exercise intensities, and strict termination criteria are warranted in individuals diagnosed with WHO Group 1 of PAH.

**Acknowledgments**

**Funding:** None.

**Footnote**

**Reporting Checklist:** The authors have completed the PRISMA reporting checklist. Available at https://dx.doi.org/10.21037/jtd-20-3296

**Conflicts of Interest:** All authors have completed the ICMJE uniform disclosure (available at https://dx.doi.org/10.21037/jtd-20-3296). The authors have no conflicts of interest to declare.

**Ethical Statement:** The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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