Prostate Cancer

Effects of Different Exercise Modalities on Fatigue in Prostate Cancer Patients Undergoing Androgen Deprivation Therapy: A Year-long Randomised Controlled Trial

Dennis R. Taaffe a,b,c,*, Robert U. Newton a,b,d,e, Nigel Spry a,f,g, David Joseph b,g,h, Suzanne K. Chambers a,i,j,k, Robert A. Gardiner a,e,l, Brad A. Wall a,m, Prue Cormie n, Kate A. Bolam c,o, Daniel A. Galvão a,b

a Exercise Medicine Research Institute, Edith Cowan University, Joondalup, WA, Australia; b School of Medical and Health Sciences, Edith Cowan University, Joondalup, WA, Australia; c School of Human Movement and Nutrition Sciences, The University of Queensland, Brisbane, QLD, Australia; d Institute of Human Performance, The University of Hong Kong, Hong Kong, China; e University of Queensland Centre for Clinical Research, University of Queensland, Brisbane, QLD, Australia; f Genesis CancerCare, Joondalup, WA, Australia; g Faculty of Medicine, University of Western Australia, Nedlands, WA, Australia; h Department of Radiation Oncology, Sir Charles Gairdner Hospital, Nedlands, WA, Australia; i Menzies Health Institute Queensland, Griffith University, Gold Coast, Australia; j Centre for Research in Cancer Control, Cancer Council Queensland, Brisbane, QLD, Australia; k Prostate Cancer Foundation of Australia, Sydney, NSW, Australia; l Department of Urology, Royal Brisbane and Women’s Hospital, Brisbane, QLD, Australia; m School of Psychology and Exercise Science, Murdoch University, Murdoch, WA, Australia; n Institute for Health & Ageing, Australian Catholic University, Melbourne, VIC, Australia; a School of Sport and Health Sciences, Stockholm, Sweden

Article info

Article history:
Accepted February 9, 2017

Associate Editor:
James Catto

Keywords:
Exercise
Fatigue
Androgen deprivation therapy
Prostate cancer
Exercise prescription
Aerobic exercise
Resistance exercise

Abstract

Background: Physical exercise mitigates fatigue during androgen deprivation therapy (ADT); however, the effects of different exercise prescriptions are unknown.

Objectives: To determine the long-term effects of different exercise modes on fatigue in prostate cancer patients undergoing ADT.

Design, setting, and participants: Between 2009 and 2012, 163 prostate cancer patients aged 43–90 y on ADT were randomised to exercise targeting the musculoskeletal system (impact loading + resistance training; ILRT; n = 58), the cardiovascular and muscular systems (aerobic + resistance training; ART; n = 54), or to usual care/delayed exercise (DEL; n = 51) for 12 mo across university-affiliated exercise clinics in Australia.

Intervention: Supervised ILRT for 12 mo, supervised ART for 6 mo followed by a 6-mo home program, and DEL received a printed booklet on exercise information for 6 mo followed by 6-mo stationary cycling exercise.

Outcome measurements and statistical analysis: Fatigue was assessed using the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire-Core 36 and vitality using the Short Form-36. Analysis of variance was used to compare outcomes for groups at 6 mo and 12 mo.

Results and limitations: Fatigue was reduced (p = 0.005) in ILRT at 6 mo and 12 mo (~5 points), and in ART (p = 0.005) and DEL (p = 0.022) at 12 mo. Similarly, vitality increased for all groups (p ≤ 0.001) at 12 mo (~4 points). Those with the highest levels of fatigue and lowest vitality improved the most with exercise (Ptrend < 0.001). A limitation was inclusion of mostly well-functioning individuals.

Conclusions: Different exercise modes have comparable effects on reducing fatigue and enhancing vitality during ADT. Patients with the highest levels of fatigue and lowest vitality had the greatest benefits.

* Corresponding author. Exercise Medicine Research Institute, School of Medical and Health Sciences, Edith Cowan University, 270 Joondalup Drive, Joondalup, WA 6027, Australia. Tel.: +61 8 63045476. E-mail address: d.taaffe@ecu.edu.au (D.R. Taaffe).

http://dx.doi.org/10.1016/j.eururo.2017.02.019
0302-2838/© 2017 European Association of Urology. Published by Elsevier B.V. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).
1. Introduction

Androgen deprivation therapy (ADT) is extensively used in the management of prostate cancer (PCa) but is associated with an array of adverse effects [1]. One adverse effect which has a considerable impact on quality of life is fatigue and a substantial proportion of men with PCa suffer from fatigue, with ~40% or more of those on long-term ADT reporting chronic fatigue or clinically-relevant fatigue which interferes with daily functioning [2].

Exercise interventions have shown positive effects on reducing or mitigating fatigue [3]. As a result, recent expert reviews in urology/oncology have incorporated aerobic and resistance exercise interventions as evidence-based strategies to mitigate toxicities from ADT including fatigue [1,4]. We [5,6] and others [7,8] have reported in relatively short-term trials (< 6 mo) of progressive resistance (strength) training and/or aerobic exercise consisting of walking/jogging or cycling at moderate to high intensity can reduce or prevent the worsening of fatigue, as can the same exercise modes when combined with dietary advice/behavioural components in a lifestyle intervention [9,10]. However, these studies have examined only the effects of short-term interventions with longer-term outcomes rarely reported. Importantly, advances to exercise protocols/prescription are required to understand the potential of different exercise modalities on fatigue. Accordingly, we report for the first time the efficacy of a 1-y long randomised controlled trial (RCT) of varying exercise interventions in PCa patients undergoing ADT with changes in fatigue and vitality assessed over 6 mo and 12 mo.

2. Patients and methods

2.1. Patients

Two-hundred and ninety-three patients with PCa were screened for participation from 2009 to September 2012 at Perth, Western Australia and Brisbane, Queensland and their progress through the study is detailed in Fig. 1. Inclusion criteria included histologically documented PCa, minimum exposure to ADT of 2 mo, without prostate-specific antigen (PSA) evidence of disease activity, and anticipated to receive ADT for the subsequent 12 mo. Exclusion criteria included bone metastatic disease, musculoskeletal, cardiovascular, or neurological conditions that could inhibit them from exercising, inability to walk 400 m or undertake exercise, and structured resistance and aerobic training in the previous 3 mo. All participants obtained medical clearance from their physician. The study was approved by the University Human Research Ethics Committee and all participants provided written informed consent.

2.2. Study design and random assignment

This was a three-armed RCT. Primary endpoints were bone mineral density and cardiovascular capacity [11], which will be reported elsewhere, with secondary endpoints including physical function and self-reported patient outcomes. Potential participants were primarily identified by their treating urologist/oncologist and referred to the study coordinator to confirm eligibility, describe the study, and obtain informed consent. Study patients underwent a familiarisation session that included correct exercise technique followed by baseline testing comprising physical tests, questionnaires, and a venous blood sample. Following baseline assessment, participants were stratified according to time on ADT (< 6 mo or ≥ 6 mo) and randomly allocated to: impact loading + resistance training (ILRT), aerobic + resistance training (ART), or to usual care/delayed exercise (DEL) by computer random assignment.

2.3. Exercise training program

ILRT was undertaken twice weekly in University-affiliated exercise clinics for 12 mo. Sessions were supervised with up to 10 participants. The impact-loading component consisted of a series of bounding, skipping, drop jumping, hopping, and leaping activities that produced ground reaction forces of 3.4–5.2 times body weight, and was progressive in nature. Specific details on progression are described elsewhere [11]. Resistance training consisted of six principal exercises that targeted the major upper and lower body muscle groups: chest press, seated row, shoulder press, leg press, leg extension, and leg curl, with supplementary exercises. Patients performed two to four sets of each exercise at an intensity of 6–12 RM (maximal weight that can be lifted 6–12 times). In addition, the ILRT group undertook home training twice weekly that consisted of two to four rotations of skipping/hopping/leaping/drop jumping [11]. ART underwent supervised exercise in the clinic twice weekly for the initial 6 mo. The aerobic-based component consisted of 20–30 min of exercise at 60–75% of estimated maximal heart rate using various modes which included walking/jogging and cycling or rowing on stationary ergometers. Resistance exercise during the initial 6 mo was the same as that undertaken in the ILRT regimen. In addition, participants were encouraged to undertake home-based aerobic activity such as walking or cycling with the goal to accumulate 150 min/wk of aerobic activity. For the 2nd 6 mo, patients were provided with a home-based maintenance program similar to our previous report [12]. DEL were provided with a printed booklet with information about exercise for the initial 6 mo, followed by 6 mo of twice weekly supervised exercise on a cycle ergometer at an intensity of ~70% maximal heart rate and flexibility exercises in the clinic. During the 12-mo study period, ILRT, ART, and DEL were asked to maintain customary physical activity and dietary patterns.

2.4. Fatigue and vitality

Study endpoints of fatigue and vitality were assessed at baseline, 6 mo, and 12 mo. Fatigue was assessed using the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire Core 30 (EORTC QLQ-C30; Table 1). Fatigue is a three-item symptom subscale with higher scores representing greater fatigue [13]. Vitality (energy level and fatigue) was assessed with the Short Form-36 Health Survey (SF-36; Table 1) [14]. The Vitality scale of the SF-36 is a four-item subdomain measure with scores ranging from 0 to 100, with higher values indicating more vitality [15].

Patient summary: We compared the effects of different exercise modes on fatigue in men on androgen deprivation therapy. All exercise programs reduced fatigue and enhanced vitality. We conclude that undertaking some form of exercise will help reduce fatigue, especially in those who are the most fatigued.
The 400-m walk was used as a measure of cardiovascular fitness [5,12,16] and muscle strength was assessed using the 1-RM method [17]. Strength is reported as the sum of the chest press and leg press, representative of upper- and lower-body strength, respectively. Percent body fat was determined using dual-energy x-ray absorptiometry. PSA and total testosterone were assessed by an accredited laboratory. Nutritional status was assessed by the Mini Nutritional Assessment [18] and self-reported physical activity by the Leisure Score Index of the Godin Leisure-Time Exercise Questionnaire [19].

2.5. Other measures

The 400-m walk was used as a measure of cardiovascular fitness [5,12,16] and muscle strength was assessed using the 1-RM method [17]. Strength is reported as the sum of the chest press and leg press, representative of upper- and lower-body strength, respectively. Percent body fat was determined using dual-energy x-ray absorptiometry. PSA and total testosterone were assessed by an accredited laboratory. Nutritional status was assessed by the Mini Nutritional Assessment [18] and self-reported physical activity by the Leisure Score Index of the Godin Leisure-Time Exercise Questionnaire [19].

2.6. Statistical analyses and sample size calculation

The sample size estimate for the RCT was based on projected changes in bone mineral density and cardiorespiratory capacity [11]. To achieve 90% power at an α level of 0.05 (two-tailed) and account for an attrition rate of up to 35%, 65 patients per group were required. For fatigue and vitality, assuming a minimally important difference (MID) of 5 points for fatigue [20] and for vitality [21], ~69 patients per group were required for fatigue (~51 patients for 80% power), and ~42 patients per group for vitality. Data were analysed using IBM SPSS Version 21 (IBM Corp., Armonk, NY, USA). Analyses included standard descriptive statistics, chi-square, one-way analysis of variance, and one-way and two-way (group x time) repeated measures analysis of variance. Follow-up tests were performed if the interaction or main effect for time was significant. Where appropriate, the Bonferroni post-hoc procedure for multiple comparisons was used to locate the source of significant differences. Trend analysis was performed using linear regression and entering quartiles of fatigue and vitality as an ordinal variable. Intention to treat was utilised for all analyses using maximum likelihood imputation of missing values (expectation maximisation). Tests were two-tailed with an α level of 0.05 applied as the criterion for statistical significance.

3. Results

3.1. Patients characteristics

There were no significant differences among groups at baseline (Table 2). The median (interquartile range) time for entry into the study since diagnosis was 8 (4–73) mo, 9 (4–47) mo, and 8 (4–40) mo, and for time on ADT 3 (2–4) mo, 3 (2–4) mo, and 2 (2–4) mo for ILRT, ART, and DEL, respectively. Of the 163 participants, four men had missing data at baseline for both fatigue and vitality resulting in a sample size of 159 men in this report. Of these 159 parti-

### Table 1 – Participant characteristics (mean ± standard deviation)

<table>
<thead>
<tr>
<th></th>
<th>ILRT (n = 57)</th>
<th>ART (n = 54)</th>
<th>DEL (n = 48)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>68.9 ± 9.1</td>
<td>69.0 ± 9.3</td>
<td>68.4 ± 9.1</td>
<td>0.947</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>173.6 ± 5.8</td>
<td>173.2 ± 6.8</td>
<td>171.6 ± 5.2</td>
<td>0.215</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>84.4 ± 11.2</td>
<td>84.9 ± 15.6</td>
<td>88.4 ± 15.4</td>
<td>0.316</td>
</tr>
<tr>
<td>Body fat (%)</td>
<td>28.1 ± 4.8</td>
<td>27.3 ± 5.9</td>
<td>29.6 ± 5.0</td>
<td>0.086</td>
</tr>
<tr>
<td>Gleason score</td>
<td>7.7 ± 1.4</td>
<td>8.0 ± 0.9</td>
<td>7.8 ± 1.0</td>
<td>0.548</td>
</tr>
<tr>
<td>Cancer stage grouping</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Localised, N (%)</td>
<td>52 (91.2)</td>
<td>50 (92.6)</td>
<td>45 (93.8)</td>
<td>0.887</td>
</tr>
<tr>
<td>Nodal metastases, N (%)</td>
<td>5 (8.8)</td>
<td>4 (7.4)</td>
<td>3 (6.3)</td>
<td></td>
</tr>
<tr>
<td>Bone metastases, N (%)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td></td>
</tr>
<tr>
<td>PSA (ng/ml)</td>
<td>1.3 ± 2.1</td>
<td>1.0 ± 1.8</td>
<td>1.3 ± 2.4</td>
<td>0.730</td>
</tr>
<tr>
<td>Testosterone (pg/ml)</td>
<td>0.8 ± 1.1</td>
<td>1.1 ± 2.6</td>
<td>1.3 ± 3.4</td>
<td>0.536</td>
</tr>
<tr>
<td>MNA</td>
<td>27.2 ± 2.3</td>
<td>27.6 ± 2.2</td>
<td>27.8 ± 1.8</td>
<td>0.633</td>
</tr>
<tr>
<td>Godin LSI</td>
<td>20.6 ± 16.5</td>
<td>23.5 ± 20.7</td>
<td>21.8 ± 16.0</td>
<td>0.698</td>
</tr>
<tr>
<td>Employed, N (%)</td>
<td>22 (38.6)</td>
<td>17 (31.5)</td>
<td>19 (39.6)</td>
<td>0.571</td>
</tr>
<tr>
<td>Married, N (%)</td>
<td>44 (77.2)</td>
<td>42 (77.8)</td>
<td>43 (89.6)</td>
<td>0.720</td>
</tr>
<tr>
<td>Current smoker, N (%)</td>
<td>3 (5.3)</td>
<td>3 (5.6)</td>
<td>3 (6.3)</td>
<td>0.822</td>
</tr>
<tr>
<td>ADT + antiandrogen, N (%)</td>
<td>27 (47.4)</td>
<td>30 (55.6)</td>
<td>27 (56.3)</td>
<td>0.586</td>
</tr>
<tr>
<td>ADT time (mo)</td>
<td>4.2 ± 4.5</td>
<td>5.3 ± 7.6</td>
<td>3.7 ± 3.7</td>
<td>0.320</td>
</tr>
<tr>
<td>Radiation, N (%)</td>
<td>49 (86.0)</td>
<td>50 (92.6)</td>
<td>40 (83.8)</td>
<td>0.341</td>
</tr>
<tr>
<td>Prostatectomy, N (%)</td>
<td>20 (35.1)</td>
<td>15 (27.8)</td>
<td>12 (25.0)</td>
<td>0.497</td>
</tr>
<tr>
<td>Other conditions</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CVD, N (%)</td>
<td>4 (7.0)</td>
<td>3 (5.6)</td>
<td>2 (4.2)</td>
<td>0.819</td>
</tr>
<tr>
<td>Hypertension, N (%)</td>
<td>20 (35.1)</td>
<td>15 (27.8)</td>
<td>23 (47.9)</td>
<td>0.104</td>
</tr>
<tr>
<td>Dyslipidaemia, N (%)</td>
<td>11 (19.3)</td>
<td>14 (25.9)</td>
<td>10 (20.8)</td>
<td>0.682</td>
</tr>
<tr>
<td>Diabetes, N (%)</td>
<td>5 (8.8)</td>
<td>7 (13.0)</td>
<td>8 (16.7)</td>
<td>0.475</td>
</tr>
</tbody>
</table>

ILRT = impact-loading + resistance training; ART = aerobic + resistance training; DEL = usual care/delayed exercise; CVD = cardiovascular disease; MNA = Mini Nutritional Assessment with malnourished <17, undernourished 17–23.5, well-nourished ≥23.5; Godin LSI, with a moderate-to-strenuous LSI ≥24 classed as active and <24 classed as insufficiently active.
of 37 men (29 in DEL) received radiation therapy in conjunction with ADT (p = 0.949). No men were on or progressed during the trial to chemotherapy. In the 1st 6-mo period, and 63% for DEL for the 6–12 mo period. Attendance at the supervised sessions was 65% and 69% for ILRT at 6 mo and 12 mo, respectively, 69% for ART for the 1st 6-mo period, and 63% for DEL for the 6–12 mo period.

### 3.2. Fatigue and vitality

There was no difference among groups for fatigue (p = 0.498) or vitality (p = 0.723) at baseline (Table 3). With training, there was no significant interaction (p = 0.304) but a significant effect for time (p < 0.001) with fatigue reduced (p = 0.005) in ILRT at 6 mo and 12 mo by ~5 points, and in ART (p = 0.005) and DEL (p = 0.022) by ~5 points at 12 mo. Similarly, there was no significant interaction (p = 0.525) but a significant effect for time (p < 0.001) with vitality increasing for all groups (p ≤ 0.001) at 12 mo by ~4 points. There was no change in fatigue or vitality during the initial 6-mo usual care period for DEL. When levels of fatigue and vitality were examined by quartiles, those with the highest levels of fatigue and lowest levels of vitality at baseline responded the best to exercise such that there was a progressive decrease in fatigue (p_{trend} < 0.001) and increase in vitality at 12 mo compared to baseline.

#### Table 3 – Fatigue and vitality at baseline, 6 mo, and 12 mo

<table>
<thead>
<tr>
<th></th>
<th>Baseline</th>
<th>6 mo</th>
<th>12 mo</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Fatigue</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ILRT</td>
<td>27.9 ± 20.7</td>
<td>22.2 ± 15.4</td>
<td>22.5 ± 16.6</td>
<td>0.005 B &gt; 6, 12</td>
</tr>
<tr>
<td>ART</td>
<td>23.4 ± 18.1</td>
<td>21.9 ± 18.4</td>
<td>17.7 ± 15.0</td>
<td>0.005 B &gt; 6, 12</td>
</tr>
<tr>
<td>DEL</td>
<td>25.8 ± 20.2</td>
<td>24.6 ± 17.7</td>
<td>20.3 ± 15.3</td>
<td>0.022 B &gt; 6, 12</td>
</tr>
<tr>
<td><strong>Vitality</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>ILRT</td>
<td>50.0 ± 10.8</td>
<td>51.9 ± 8.0</td>
<td>54.6 ± 8.5</td>
<td>&lt;0.001 B, 6 &lt; 12</td>
</tr>
<tr>
<td>ART</td>
<td>51.5 ± 10.7</td>
<td>52.7 ± 9.8</td>
<td>55.3 ± 8.7</td>
<td>0.001 B, 6 &lt; 12</td>
</tr>
<tr>
<td>DEL</td>
<td>50.3 ± 10.0</td>
<td>50.1 ± 9.7</td>
<td>53.9 ± 8.1</td>
<td>&lt;0.001 B, 6 &lt; 12</td>
</tr>
</tbody>
</table>

*ART = aerobic + resistance training; B = baseline; DEL = usual care/delayed exercise; ILRT, impact-loading + resistance training.*
in vitality ($p_{\text{trend}} < 0.001$) with exercise (Fig. 2). Sensitivity analysis was conducted using complete cases [22] with no change in the results for vitality and although for fatigue there was a significant main effect for time ($p = 0.003$) with the magnitude of difference within groups over time similar, with the reduction in sample size and hence statistical power the only significant difference was for ART ($p = 0.032$).

### 3.3. Other measures and adverse events

For cardiorespiratory fitness, there was no significant interaction ($p = 0.216$) but a significant effect for time ($p < 0.001$) with fitness improved ($p = 0.008$) in ILRT at 12 mo by $\sim 14$ s and in ART ($p < 0.001$) by $\sim 13$ s at 12 mo, with the change in DEL of $\sim 1.1$ s at 12 mo approaching significance ($p = 0.063$; Table 4). Notably, there was no change in cardiorespiratory fitness during the nonexercise period for DEL. For muscle strength, there was a significant interaction ($p < 0.001$) with strength progressively increasing at 6 mo and 12 mo ($p < 0.001$) in ILRT, increasing during the initial 6-mo supervised phase in ART ($p < 0.001$) with no change thereafter, and no change between baseline and 6 mo in DEL but a difference by 12 mo following exercise ($p < 0.001$). No adverse effects from exercise resulted in any participants having to withdraw from the study. Two men in ILRT withdrew within the 1st 6 mo due to compressed spinal discs and shoulder issues, although the person developed shoulder issues prior to commencing exercise training. Two men in ART had cardiovascular problems, one in the 1st 6 mo and one in the 2nd 6 mo, with one requiring heart bypass surgery while another participant in ART developed back pain. Two men withdrew from DEL in the 1st 6 mo due to difficulty walking and the other required back surgery.

### 4. Discussion

To our knowledge, this is the first year-long RCT to evaluate the effects of different exercise modalities on fatigue in prostate cancer patients undergoing ADT. There were two important findings: (1) all exercise modalities (resistance + impact loading, aerobic + resistance, aerobic only) had a beneficial effect on fatigue and vitality, and (2) those with the highest levels of fatigue and lowest vitality improved the most with exercise.

A substantial proportion of PCa patients will receive ADT as part of their treatment. Prevalence of clinically-relevant fatigue in these patients has been reported as $\sim 40\%$ [22,23]. The fatigue scores from the EORTC QLQ-C30 in our cohort are similar to the reference values for men with PCa aged 60–69 y (25.2 $\pm$ 26.6) [24]. Vitality has also been shown to decline during ADT. In a population-based sample of men with PCa followed for 1 y, Alibhai et al [25] reported vitality was substantially lower in ADT-treated compared with non-ADT patients. A more recent prospective study also showed changes in vitality following 12 mo of ADT with rapid declines during the initial 3 mo [26].

Short-term exercise trials have shown the efficacy of combined resistance and aerobic exercise, or as sole training modes, to improve fatigue in men undergoing radiation therapy/ADT for PCa [5,7,8]. We have also shown improvements in vitality following a 12-wk exercise intervention [5]. Here we extend these findings by providing data on the largest exercise trial undertaken with men undergoing ADT.
for PCa by examining the long-term effects of exercise and potential impact of different exercise modalities. We found that all exercise modalities had a similar effect on fatigue and vitality following the intervention. In a recent systematic review on fatigue instruments, the MID reported for the EORTC QLQ-C30 ranged from 3.0–19.7 points [20]. The mean differences for our exercise regimens was ~5 points with those in quartiles 3 and 4 at baseline having a change of 10 or more points. Similarly, it has been recommended that the MID in vitality using the SF-36 is 5 points [21]. The mean change for each of the three exercise regimens was ~4 points, with quartiles 3 and 4 gaining a mean of 5 points or more. This result is important as it provides practical information to guide the prescription of exercise in men with PCa to mitigate cancer-related fatigue.

We have recently reported that only ~12% of Australian PCa survivors are meeting sufficient exercise levels [27]. It appears that supervised exercise undertaken at moderate- to high-intensity, irrespective of modality (eg, aerobic, resistance, or impact) has a beneficial effect on fatigue. Moreover, those with higher levels of fatigue/lower levels of vitality responded the best to exercise such that there was a progressive decrease in fatigue and increase in vitality with exercise. As a result, fatigued patients are likely to benefit most from any form of structured supervised exercise when undertaken at appropriate intensity and dose. From this we propose that screening patients on ADT for fatigue and directing tailored and prescribed exercise interventions to these men should be part of the prostate cancer care pathway.

During the nonexercise period for the delay group there was no change in fatigue or vitality and no change in cardiorespiratory fitness or muscle strength. Conversely, with exercise improvements were observed in physical functioning as they were in ILRT and ART, and these were accompanied by changes in fatigue and vitality. Cardiorespiratory fitness changes as determined by the 400-m walk [28], although not substantial [16], would at least provide a greater safety margin before thresholds for disability are encountered, hence may potentially be clinically meaningful (especially for men in poorer condition than those in the present study).

Our study has several features that are worthy of comment. This is the largest and longest exercise trial in PCa patients undergoing ADT examining different exercise modalities including resistance, impact loading and aerobic training. Fatigue was assessed using the EORTC QLQ-C30 fatigue subscale which is a validated measure, widely used, and recommended for use in trials to measure cancer-related fatigue [29]. However, the generalisability of the data may be limited given that participants volunteered to participate in an exercise trial, were generally quite healthy, and predominantly were married and nonsmokers. In addition, a potential confounding factor was the group nature of the supervised sessions resulting in the sharing of common experiences and the camaraderie which may have developed, impacting on the outcomes of the study. Finally, men in this study were primarily in the initial year of ADT, therefore results may not be generalisable to men receiving ADT for a longer duration.

5. Conclusions

In conclusion, in the largest year-long exercise intervention study in men with PCa undergoing ADT, all exercise programs had comparable effects on reducing fatigue and enhancing vitality. However, the benefits were small to nonexistent for those least fatigued at baseline and as such an involved intervention should primarily be considered for those who are most fatigued. Encouraging fatigued patients to undertake exercise at adequate intensity, regardless of mode, will likely aid in reducing or attenuating the adverse effects of ADT on fatigue and vitality. Screening all men on ADT for fatigue and providing an exercise intervention is warranted.

Author contributions: Dennis R. Taaffe had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

Study concept and design: Taaffe, Newton, Spry, Joseph, Galvão.
Acquisition of data: Taaffe, Newton, Wall, Bolam, Galvão.
Analysis and interpretation of data: Taaffe, Newton, Spry, Galvão.
Drafting of the manuscript: Taaffe, Newton, Spyr, Galvão.
Critical review of the manuscript for important intellectual content: Taaffe, Newton, Spry, Joseph, Gardiner, Bolam, Wall, Cormie, Galvão.
Statistical analysis: Taaffe, Newton, Galvão.
Obtaining funding: Taaffe, Newton, Spry, Joseph, Gardiner, Galvão.
Administrative, technical, or material support: None.
Supervision: None.
Other: None.

Financial disclosures: Dennis R. Taaffe certifies that all conflicts of interest, including specific financial interests and relationships and affiliations relevant to the subject matter or materials discussed in the manuscript (eg, employment/affiliation, grants or funding, consultancies, honoraria, stock ownership or options, expert testimony, royalties, or patents filed, received, or pending), are the following: None.

Funding/Support and role of the sponsor: This study was funded by the National Health and Medical Research Council 534409, Prostate Cancer Foundation of Australia, Cancer Council of Western Australia, and Cancer Council of Queensland. The sponsors did not participate in the design or conduct of the study; collection, management, analysis, and interpretation of the data; or in the preparation, review, or approval of the manuscript. Daniel A. Galvão is funded by a Cancer Council Western Australia Research Fellowship. Suzanne Chambers is supported by an Australian Research Council Professorial Future Fellowship.

A phase 3 clinical trial of exercise modalities on treatment side-effects in men receiving therapy for prostate cancer; ACTRN12609000200280

References


