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Vigorous intensity aerobic interval exercise in bladder cancer patients prior to radical cystectomy: a feasibility randomized controlled trial

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ABSTRACT

**Purpose:** Strategies to improve pre-operative cardiopulmonary fitness could positively impact recovery after surgery. This study investigated the feasibility of vigorous intensity aerobic interval exercise in bladder cancer patients prior to radical cystectomy (RC). **Methods:** A total of 60 patients were randomised (1:1) to exercise or control following a cardiopulmonary exercise test (CPET). The exercise group was offered twice-weekly pre-operative supervised vigorous intensity aerobic interval exercise in addition to standard treatment. The controls received standard treatment only. A repeat CPET was undertaken before surgery and post-operative recovery outcomes were recorded. **Results:** Over half of the 112 eligible patients approached in the clinic were recruited to the study (53.5%), with recruited patients attending a median of 8 (range: 1-10) exercise sessions over a pre-operative period of 3-6 weeks. Improvements in peak values of oxygen pulse \( (P=0.001) \), minute ventilation \( (P=0.002) \) and power output \( (P<0.001) \) were observed at the follow-up CPET in the exercise group versus controls and there were no adverse events. Although this feasibility study was not powered to detect changes in post-operative recovery outcomes, there were marginal (non-significant) differences in favour of the exercise group in post-operative Clavien-Dindo score and need for High Dependency Unit inotropic support.

**Conclusions:** Bladder cancer patients respond well to pre-surgical aerobic interval exercise and the improvements in cardiopulmonary fitness variables could have important implications for post-operative recuperation after RC. These findings provide a strong foundation for an adequately powered randomized controlled trial.

**Key words:**

Pre-operative care, exercise, urinary bladder neoplasms
INTRODUCTION

Pre-operative cardiopulmonary fitness is increasingly being recognised as an important factor influencing post-operative recovery outcomes. Almost two decades ago, Older et al. identified an association between low cardiopulmonary fitness in older people and poor outcome following major surgery [1]. Other studies have since provided further evidence of the utility of pre-operative cardiopulmonary exercise test (CPET) variables for risk stratification of surgical patients, including those undergoing radical cystectomy (RC) [2, 3]. However, the role of pre-operative exercise programs (prehabilitation) for optimising cardiopulmonary fitness prior to surgery has received less attention [4]. Although systematic review evidence suggests that exercise training can improve cardiopulmonary fitness before surgery in older people and can reduce the risk of post-operative complications following major abdominal surgery [5-7], the paucity of high quality clinical trials has been highlighted [5]. Furthermore, studies that have investigated the effects of exercise prehabilitation programs on post-operative recovery outcomes cancer patients have yielded equivocal results.

Intensive supervised pre-operative exercise programs resulted in modest improvements in pre-operative cardiopulmonary fitness measures in lung cancer patients, including the six-minute walk test [8] and peak aerobic capacity [8, 9], but with no effect on pre-surgical or longer-term quality of life [10]. In addition, a supervised exercise program involving trunk and limb-strengthening exercise improved chemoradiotherapy completion rate in patients with gastrointestinal cancers, but did not improve post-operative recovery outcomes [11]. In contrast, Sekine et al. [12] reported a reduction in post-operative pulmonary complications and hospital length of stay in lung cancer patients with chronic obstructive pulmonary disease after daily hospital-based pulmonary exercises and walking (5000 steps/day) in the two weeks prior to lobectomy. Similarly, in prostate cancer patients undergoing radical prostatectomy, there is evidence that home-based pelvic floor muscle exercise has favourable effects on post-operative urinary continence outcomes up to 12 months of follow-up [13-15]. However, home-based exercise programs involving aerobic and resistance exercise have had minimal impact on post-operative hospital length of stay or severity of complications in colorectal or bladder cancer patients [16-19], although Jensen et al. [19] reported improved post-operative mobility in the latter.

Because of the short time-window between decision for surgery and RC, there is a need to optimise the exercise stimulus for cardiopulmonary adaptations and the potential advantages of vigorous intensity interval exercise in this respect, were recently highlighted [20]. Interval training enables patients to undertake aerobic (endurance) exercise at a higher intensity than would be possible for continuous exercise at the same intensity but the
feasibility of this exercise modality in bladder cancer patients awaiting RC is unknown. Hence, the main purpose of this study was to investigate the feasibility of randomizing bladder cancer patients to a short-term program of pre-operative vigorous intensity aerobic interval exercise versus standard care prior to elective RC. A secondary aim was to collect and report preliminary data on CPET and post-operative recovery outcomes before and after the exercise prehabilitation program.

METHODS

Patient recruitment, randomization and sample size

Patients were recruited from the Urology Department at the Norfolk and Norwich University Hospitals National Health Service (NHS) Foundation Trust, United Kingdom, between 2012 and 2014, and were randomized to standard treatment or pre-operative exercise training. Medical clearance to participate in the study was given by one of four consultant urologists, who were blind to treatment allocation. Randomization was undertaken using a pre-generated random sequence (nQuery Advisor 6.1; Statistical Solutions) which was held by a research administrator not involved in the day to day running of the study. Upon completion of the baseline assessments, the study urologist (SB) phoned the administrator for details of group allocation (exercise or standard care). As the main purpose was to assess the feasibility of the exercise program, there was no formal sample size calculation but we aimed to recruit 60 patients [30 in each group) in accordance with published recommendations [21, 22]. Written informed consent was obtained prior to study participation and ethical approval was granted by the East of England Regional Ethics Committee.

Inclusion and exclusion criteria

Bladder cancer patients listed for RC via the multi-disciplinary team were included in the study (eligibility was not limited by choice of surgical technique). Patients undergoing neo-adjuvant chemotherapy before RC were also included but entered the trial at least 2 weeks after completion of treatment. Patients offered urinary diversion for benign disease were excluded from the study.

Standard treatment

Patients are treated according to targets set by NHS England, i.e. following general practitioner referral for a suspected cancer, patients are to be investigated by 31 days and treated by 62 days, resulting in a time-window between decision to operate and RC of 31 days (though this can vary due to medical decisions and availability
of operating slots). Pre- and post-operatively, patients in both groups received the same level of standard

treatment, which included pre-planned admission to the same High Dependency Unit (HDU) in the immediate
post-operative period and subsequent step down to a urology ward. Patients were cared for by the same group of
nurses and doctors who were blinded to group allocation. Patients in the control group were advised to carry on
with their lifestyles in the ‘usual way’.

Supervised exercise intervention

Twice-weekly exercise training sessions were supervised by a team of exercise science staff working closely
with the study urologist (SB) in an exercise facility at the University of East Anglia, UK, which is close to the
treating hospital. Sessions comprised vigorous intensity aerobic interval exercise on a cycle ergometer (Monark
824E; Varberg, Sweden) using the Borg Ratings of Perceived Exertion (RPE) Scale to control intensity [23].
The Borg RPE Scale is a psychophysiological scale, ranging from “No exertion” to “Maximal Exertion”.
Following a 5-10 min warm-up against light resistance (50 W), the aim was for patients to perform 6 x 5 min
intervals at a target perceived exertion of 13-15 (‘somewhat hard’ to ‘hard’, equating to 70-85% predicted
maximum heart rate based on 220-age), with 2.5 min interpolated active rest intervals against light resistance
(50 W). They were instructed to maintain a steady pedalling cadence of 50-60 rev-min⁻¹ during the aerobic
intervals and the exercise program was progressed by gradually adding more load to the flywheel to maintain
the target perceived exertion (Figure 2). Immediately following the aerobic intervals, patients performed a ‘cool-
down’ against low resistance (50 W).

Feasibility outcomes

Feasibility was assessed in terms of recruitment and attrition, willingness to be randomized, acceptability of the
outcome measures, adherence to the intervention, safety and suitability of the exercise dose and adverse events.

Cardiopulmonary exercise test and post-operative recovery outcomes

Cardiopulmonary exercise tests (CPETs) and post-operative recovery outcomes were blindly assessed by an
exercise physiologist who was not directly involved in the supervision of exercise sessions and clinical staff at
the treating hospital who were unaware of group allocation, respectively.
All patients performed a baseline incremental CPET to maximum exercise tolerance on an electronically-braked cycle ergometer (Excalibur Sport, Lode, Netherlands). Following a 2-min warm-up against no resistance (0 W), work rate was increased using a ramp protocol (10-20 W/min) to maximum exercise tolerance. Heart rate was recorded continuously by ECG (Cardioperfect, Cardioperfect, Welch Allyn, USA). Pulmonary gas exchange variables (\(\dot{V}O_2\), \(\dot{V}CO_2\), minute ventilation \([\dot{V}E]\) and other respiratory variables) were measured breath-by-breath with an on-line expired gas analysis system (Ultima, CardioO2; Medical Graphics Corporation). AT was determined from 30 s averaged data by two experienced exercise physiologists (JS and GC) using the V-slope method and confirmed by analysing the ventilatory equivalents [24]. Peak values for all variables were recorded as the highest value over any 30-s averaged period, with peak oxygen pulse calculated as the amount of oxygen consumed per heartbeat.

**Post-operative recovery outcomes**

Clavien-Dindo grading was used to score post-surgical complications [25, 26]. Post-operative ileus and chest infection prevalence, time spent in HDU, need for inotropic support and hospital length of stay (LoS) were used as other post-operative recovery outcomes.

**Data analysis**

All statistical analyses were undertaken using SPSS (SPSS Inc., Chicago, Illinois, USA). CPET and post-operative recovery outcomes were tested for normality of distribution using Kolmogorov-Smirnov test. BMI and CPET variables were normally distributed and analysed using ANCOVA, with baseline values as the covariate [27], to compare differences between the groups at follow-up (prior to surgery). Post-operative recovery outcomes were not normally distributed and non-parametric statistical tests were used. The Chi-Square test was used to compare categorical data. The Spearman’s rank correlation coefficient was used to investigate bivariate associations between CPET variables and post-operative recovery outcomes in pooled baseline data (independent of group assignment). Normally distributed data are presented as mean ± SD and non-normally distributed data as medians and range (unless otherwise stated). The significance level was set at \(P<0.05\).
RESULTS

Feasibility

Of the 112 eligible patients who were approached to take part in the study, 60 (53.5%) agreed to participate (53 males; 7 females). All recruited patients were willing to be randomized and none withdrew consent following randomization. The groups were well-matched for demographic variables, surgical treatment, comorbidities and smoking habits, though more patients in the control group had a history of ischemic heart disease and more patients in the exercise group had undergone neoadjuvant chemotherapy prior to the study (Table 1). Travel distance to the exercise facility was cited as the reason for non-participation in 38 (73%) of 52 eligible patients who declined to take part. Three eligible patients (5.8%) refused to take part because they felt they would be unable to undertake the CPET or comply with the exercise regimen. A further four patients (7.6%) refused to take part as they had an indwelling urethral catheter or a nephrostomy which they thought might prevent them from undertaking exercise. Seven patients (13.4%) did not specify their reasons for not participating. Only five of the 60 recruited patients dropped out of the study, two were deemed unfit for surgery following randomization and three opted for radiotherapy after the follow-up CPET. Recruitment and patient flow through the study is shown in Figure 1.

Most patients completed all assessments, demonstrating the acceptability of the pre-operative CPET and post-operative recovery outcomes. One patient from each group dropped out of the study before the follow-up CPET. Another two patients from the exercise group and four controls did not complete the follow-up CPET but their post-operative outcomes were included in the analysis (Figure 1). This means that a total of 27/30 (90%) of patients in the exercise group and 25/30 (83%) patients in the control group completed both CPET assessments (Figure 1), with no adverse events and with AT data being available for 24 and 23 patients, respectively. Post-operative recovery outcomes were available for 27/30 (90%) patients in the exercise group and 28/30 (93%) patients in the control group (Figure 1).

The median number of supervised exercise sessions attended by patients in the exercise arm was 8 (range: 1-10) over a pre-operative period of 3 - 6 weeks. Three exercise patients underwent RC <4 weeks after recruitment and consequently attended <8 sessions. Between the first and fourth week, patients exercised at an average of 85-87% predicted maximum heart rate (based on 220-age) and 90-92% of the measured peak HR during the CPET (Figure 2), demonstrating the safety and suitability of the exercise dose. Flywheel load (power output)
was gradually increased from $111 \pm 5.5 \, W$ to $122 \pm 5.8 \, W$ during this time to ensure target HRs for vigorous intensity exercise were maintained as patients adapted to the exercise program (Figure 2). The average number of aerobic intervals achieved in the first week of exercise was 5.5 (range: 3.5-6.0), whereas all patients were achieving 6 intervals per session in the fourth week. There were no adverse events resulting from the supervised exercise sessions.

**CPET and post-operative recovery outcomes**

CPET variables at baseline and follow-up are presented in Table 2. The time between baseline and follow-up CPET was 32 (6.5) days (range: 20-45 days) for the exercise group and 29 (4.2) days (range: 21-37 days) for the controls. Improvements in peak values of oxygen pulse ($P=0.001$), $\bar{V}E$ ($P=0.002$) and power output ($P<0.001$) but not peak $\bar{V}O_2$ ($P=0.057$) or AT ($P=0.637$) were observed at the follow-up CPET in the exercise group versus controls (Table 2). Although this feasibility study was not powered to detect changes in post-operative recovery outcomes, four patients (15%) in the exercise group versus 10 controls (36%) had some deviation from the normal post-operative course of recovery (Clavien-Dindo Grade ≥1; $P=0.075$), whereas one patient (4%) in the exercise group and four controls (14%) had a Clavien-Dindo Grade ≥ 3; ($P=0.172$). The prevalence of post-operative ileus ($P=0.808$) and chest infection ($P=0.609$) were similar between the exercise and control groups (6 versus 7 patients and 3 versus 2 patients, respectively). Median time spent in HDU was 1 day in both the exercise and control group (range 1-10 and 1-7 days, respectively; $P=0.938$), although fewer patients in the exercise group needed HDU inotropic support, reaching borderline statistical significance (2 versus 7 patients; $P=0.078$). The median hospital LoS was 7 days in both the exercise and control groups (range 4-78 and 5-107 days, respectively; $P=0.865$). When the pre-operative data were pooled and considered independent of group assignment, inverse associations were observed between LoS (HDU and hospital) and pre-operative CPET variables ($P<0.05$; Table 3).

**DISCUSSION**

This is the first study to show that vigorous intensity aerobic interval exercise is feasible in bladder cancer patients awaiting RC. Patients adapted to the program quickly and all were able to perform six aerobic intervals per exercise session at the vigorous intensity target HR by the fourth week of exercise. The demographic profile of our patients was representative of bladder cancer patients undergoing RC in terms of age, comorbidities and risk factors, and the recruitment rate of eligible patients was very good (53.5%). The main reason for non-
participation amongst eligible patients was distance between the exercise facility and their home. As the hospital serves a large rural community, with some patients living >80 km away, twice-weekly travel to the supervised exercise sessions was considered to be too onerous by some participants. We also observed excellent retention over the course of the study (8.3% attrition), which is likely to be at least in part due to flexible exercise scheduling (i.e. exercise sessions were arranged at suitable times for patients). In addition, most patients (87%) were able to attend both CPET assessments prior to RC and there were no adverse events. The time between baseline and follow-up CPET assessments was in the range of 20-45 days across both groups. This provided an opportunity for all patients in the exercise arm to attend a minimum of six supervised exercise sessions prior to RC.

Because of the short time-window between decision for surgery and RC, there is a need for pre-surgical exercise programs to be effective and time efficient and the potential advantages of vigorous interval exercise were recently highlighted [20]. Our exercise program involved aerobic interval exercise training at “somewhat hard” to “hard” intensity (Borg RPE 13-15), equating to ~70-85% predicted maximum heart rate [28], which is reported to be optimal for inducing improvements in cardiopulmonary fitness in previously sedentary older people [29]. Improvements in peak values of oxygen pulse, $\dot{V}E$ and power output in the exercise group suggests that the exercise program could be an effective stimulus for inducing cardiopulmonary adaptations in the short time available before surgery. The improvement in peak oxygen pulse and progressive increase in flywheel load (power output) needed to maintain the vigorous intensity target HR (Figure 2) are indicative of adaptations in cardiac stroke volume, which commonly result from endurance training [30]. These improvements were accompanied by a marginal (non-significant) increase in peak $\dot{V}O_2$, which in conjunction with enhanced pedalling efficiency following exercise training, probably account for the increase in peak power output observed in the exercise group. In contrast, AT was unchanged at the CPET follow-up, suggesting metabolic adaptations influencing oxygen utilisation within the active skeletal muscles were negligible. In the exercise group, the ability of skeletal muscle to utilise any additional oxygen resulting from enhanced delivery may have been compromised to a greater degree because a higher proportion of participants had undergone neoadjuvant chemotherapy prior to the study (33% versus 17% in the control group). Neoadjuvant chemotherapy was previously shown to adversely affect skeletal muscle mitochondrial oxidative phosphorylation in patients with locally advanced rectal cancer, probably influencing the reported decline in peak $\dot{V}O_2$ and AT prior to surgery [31]. Furthermore, cisplatin (commonly used in the neoadjuvant chemotherapeutic treatment of bladder cancer
patients) causes skeletal muscle atrophy and alterations in a range of metabolic signalling pathways that were only partially reversed by exercise training in healthy mice [32]. As improvements in AT following endurance training are strongly influenced by adaptations in metabolic pathways that enhance fat utilization and oxidative ATP production [30], a higher volume of pre-operative aerobic interval exercise and/or longer period of recovery may be needed to overcome the adverse effects of neoadjuvant chemotherapy on skeletal muscle mitochondria and signalling pathways prior to RC. In this respect, a 6-week program of thrice-weekly cycle ergometer interval training was needed to restore AT to pre-neoadjuvant chemotherapy levels in patients with locally advanced rectal cancer [33]. Given the feasibility of twice-weekly vigorous aerobic interval exercise in this patient group, programs which offer additional weekly sessions could be tested and might enhance the stimulus for metabolic adaptations, particularly in patients who have recently undergone neoadjuvant chemotherapy.

Systematic reviews have reported encouraging evidence supporting the positive impact of pre-operative exercise on post-operative recovery outcomes in patients undergoing cardiac and abdominal surgery but there has been significant heterogeneity in study design and a paucity of high quality evidence [34, 35]. Our results for post-operative recovery outcomes are to be considered with caution as adequately powered clinical trials are needed to establish whether improvements in pre-operative cardiopulmonary fitness are associated with enhanced recovery following RC. Nevertheless, these preliminary data show marginal (non-significant) differences in favour of the exercise group in post-operative Clavien-Dindo score (≥1) and need for HDU inotropic support. It has been proposed that pre-operative exercise has the potential to improve physiological reserve, thereby helping to counter the physical challenges of surgery and enhancing post-operative recuperation [4]. However, the magnitude of change in pre-operative cardiopulmonary fitness required to positively influence surgical outcomes is unclear. Previous research suggests that improvements of 2-3 ml kg⁻¹ min⁻¹ in pre-operative peak \( \dot{V}O_2 \) following supervised aerobic exercise programs [8, 9] might not translate into quality of life enhancements in lung cancer patients before or after surgery [10]. Other exercise prehabilitation studies in lung, colorectal and bladder cancer patients have not reported changes in cardiopulmonary fitness following exercise prehabilitation programs that resulted in improvements [12] or no improvements [16-19] in post-operative recovery outcomes. Observational studies suggest that an AT of less than 11-12 ml kg⁻¹ min⁻¹ is associated with major post-operative complications (Clavien Class ≥3) and poorer recovery outcomes (including hospital length of stay) in bladder cancer patients undergoing RC [2, 36] and this is consistent with evidence from a large-scale study of
patients (N=843) undergoing intra-abdominal surgery [37]. AT was >11 ml·kg⁻¹·min⁻¹ in >50% of our participants prior to surgery and when our data were pooled and considered independent of group assignment, higher scores for pre-operative peak $\dot{V}O_2$, peak $\dot{V}E$, peak power output and AT were associated with improved recovery outcomes (Table 3). This observational evidence, in conjunction with the lack of intervention studies linking changes in pre-operative cardiopulmonary fitness to post-operative recovery outcomes, raises the question of whether cardiopulmonary fitness gains that fail to reach a given AT or peak $\dot{V}O_2$ threshold could positively influence post-operative recovery outcomes in this patient group. In addition, the relative importance of cardiopulmonary fitness versus other factors in optimising post-operative recovery, e.g. adequate nutrition, mental preparation, etc. [4], is an important avenue for future research.

This study had some limitations. Firstly, the sample size was insufficient to draw definitive conclusions about the effects of improved pre-operative cardiopulmonary fitness on post-operative recovery. Although we showed that the exercise program has the potential to improve indices of cardiopulmonary fitness, a larger-scale randomized controlled trial with a sufficient number of patients to detect changes in key recovery outcomes and longer-term quality of life, is needed to address this important question. Secondly, as patients were recruited from a single centre serving a large rural community, the recruitment rate and reasons for non-participation may be less representative of patients living closer to treating hospitals. Travel distance to the exercise facility was cited as the main reason for non-participation in the study but this is likely to be less of an issue in patients living in closer proximity. Nevertheless, other reasons (including physical limitations or a lack of confidence to engage in the exercise program) prevented >26% of eligible patients from taking part. This suggests that a program of hospital supervised vigorous intensity aerobic interval exercise is unlikely to be feasible for all bladder cancer patients. For this reason, other forms of pre-operative exercise, particularly exercise that can be undertaken in the home or community environment, should be evaluated in future research.

In conclusion, the results of this study show that vigorous intensity aerobic interval exercise in the short time-window between decision to operate and RC was feasible in a significant proportion (54%) of bladder cancer patients being treated in a large Urology Department. In addition, we present preliminary evidence of improvements in some cardiopulmonary fitness variables that could potentially impact on post-operative recovery outcomes. Because of the small sample size, it is not possible to make definitive conclusions about the impact of pre-operative improvements in cardiopulmonary fitness on post-operative recovery outcomes.
Nevertheless, these encouraging preliminary data provide a strong foundation for an adequately powered randomized controlled trial.

**CONFLICT OF INTEREST STATEMENT**

All authors declare that they have no conflicts of interest. The corresponding author has full control of all primary data and agrees to allow the journal to review the data if requested.

**REFERENCES**


FIGURE LEGENDS

**Figure 1.** Participant flow through the trial.

**Figure 2.** Average weekly heart rate (HR; top figure) and power output (bottom figure) over four pre-operative weeks of supervised aerobic interval exercise. Average heart rate data are presented as percentage of the peak heart rate measured during the pre-operative CPET and predicted maximum heart rate based on 220 – age.
Figure 1. Participant flow through the trial
Figure

Weeks of supervised exercise

% maximum HR

% Measured peak HR

% Predicted maximum HR

Average power output (W)

Weeks of supervised exercise
Table 1. Baseline characteristics of participants allocated to exercise plus standard care or standard care only. Data are presented as mean ± SD or numbers (percentages).

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Exercise group (N=30)</th>
<th>Control group (N=30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (range)</td>
<td>71.60 ± 6.80</td>
<td>72.5 ± 8.40</td>
</tr>
<tr>
<td>Female</td>
<td>3 (10)</td>
<td>4 (13)</td>
</tr>
<tr>
<td>White</td>
<td>30 (100)</td>
<td>30 (100)</td>
</tr>
<tr>
<td><strong>Anthropometric variables and blood pressure</strong></td>
<td></td>
<td></td>
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<tr>
<td>Height (m)</td>
<td>1.73 ± 6.85</td>
<td>1.68 ± 8.05</td>
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<tr>
<td>Body mass (Kg)</td>
<td>81.17 ± 13.38</td>
<td>76.20 ± 11.60</td>
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<tr>
<td>BMI (kg·m⁻²)</td>
<td>27.09 ± 4.20</td>
<td>26.91 ± 4.45</td>
</tr>
<tr>
<td><strong>Comorbid conditions</strong></td>
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<td></td>
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<tr>
<td>Hypertension</td>
<td>17 (57)</td>
<td>17 (57)</td>
</tr>
<tr>
<td>Ischemic heart disease</td>
<td>3 (10)</td>
<td>8 (27)</td>
</tr>
<tr>
<td>Type 2 diabetes</td>
<td>3 (10)</td>
<td>4 (13)</td>
</tr>
<tr>
<td><strong>Bladder cancer treatment</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Neoadjuvant chemotherapy</td>
<td>10 (33)</td>
<td>5 (17)</td>
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<tr>
<td>Surgical treatment</td>
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<tr>
<td>Laparoscopic</td>
<td>27 (90)</td>
<td>28 (93)</td>
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<tr>
<td>Open</td>
<td>1 (3)</td>
<td>1 (3)</td>
</tr>
<tr>
<td>Neo bladder formation</td>
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<td>1 (3)</td>
</tr>
<tr>
<td><strong>Smoking status</strong></td>
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<tr>
<td>Current smoker</td>
<td>4 (13)</td>
<td>4 (13)</td>
</tr>
<tr>
<td>Ex-smoker</td>
<td>20 (66)</td>
<td>13 (43)</td>
</tr>
<tr>
<td>Never smoker</td>
<td>6 (20)</td>
<td>13 (43)</td>
</tr>
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</table>
Table 2. Cardiopulmonary variables at baseline and follow-up (prior to radical cystectomy). Data are presented as mean ± SD, with 95% confidence intervals in parentheses.

<table>
<thead>
<tr>
<th>CPET variables</th>
<th>Control group</th>
<th>Exercise group</th>
<th>Adjusted mean difference at follow-up CPET</th>
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<tbody>
<tr>
<td></td>
<td>Baseline</td>
<td>Pre-surgery</td>
<td>Baseline</td>
<td></td>
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<tr>
<td>Peak $\dot{V}O_2$ (ml.kg$^{-1}$.min$^{-1}$)</td>
<td>20.38 ± 5.59 (18.07-22.68)</td>
<td>20.84 ± 5.43 (18.60-23.08)</td>
<td>19.22 ± 4.80 (17.32-21.12)</td>
<td>21.07 ± 5.60 (18.85-23.29)</td>
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<td>PPO (W)</td>
<td>131 ± 36 (116-145)</td>
<td>129 ± 44 (111-147)</td>
<td>131 ± 39 (116-146)</td>
<td>148 ± 41 (132-165)</td>
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<tr>
<td>Peak HR (beats.min$^{-1}$)</td>
<td>143 ± 24 (134-153)</td>
<td>143 ± 23 (134-152)</td>
<td>139 ± 25 (129-149)</td>
<td>137 ± 26 (127-148)</td>
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<tr>
<td>Peak RER</td>
<td>1.38 ± 0.13 (1.32-1.43)</td>
<td>1.35 ± 0.12 (1.30-1.40)</td>
<td>1.36 ± 0.11 (1.32-1.40)</td>
<td>1.35 ± 0.12 (1.30-1.40)</td>
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<tr>
<td>Peak $\dot{V}E$: (L.min$^{-1}$)</td>
<td>67.04 ± 19.50 (58.99-75.09)</td>
<td>68.07 ± 19.30 (60.10-76.03)</td>
<td>70.33 ± 22.54 (61.41-79.25)</td>
<td>78.63 ± 23.12 (69.48-87.78)</td>
</tr>
<tr>
<td>Peak OP (ml.beat$^{-1}$)</td>
<td>10.72 ± 2.24 (9.79-11.64)</td>
<td>10.83 ± 2.33 (9.87-11.79)</td>
<td>11.31 ± 2.74 (10.23-12.40)</td>
<td>12.74 ± 2.88 (11.60-13.88)</td>
</tr>
<tr>
<td>AT (ml.kg$^{-1}$.min$^{-1}$)</td>
<td>11.38 ± 2.57 (10.27-12.49)</td>
<td>12.21 ± 2.63 (11.07-13.35)</td>
<td>11.49 ± 2.08 (10.61-12.37)</td>
<td>12.00 ± 2.97 (10.74-13.25)</td>
</tr>
<tr>
<td>OP at AT (ml.beat$^{-1}$)</td>
<td>8.33 ± 2.05 (7.45-9.22)</td>
<td>8.56 ± 1.99 (7.70-9.42)</td>
<td>8.90 ± 2.06 (8.03-9.77)</td>
<td>9.76 ± 2.63 (8.65-10.87)</td>
</tr>
<tr>
<td>$\dot{VE}/\dot{V}CO_2$ at AT</td>
<td>31.00 ± 5.09 (28.80-33.20)</td>
<td>31.17 ± 4.95 (29.03-33.31)</td>
<td>31.50 ± 4.11 (29.77-33.23)</td>
<td>31.17 ± 3.27 (29.79-32.55)</td>
</tr>
</tbody>
</table>

$\dot{V}O_2$: oxygen consumption per minute; PPO: peak power output; HR: heart rate; RER: respiratory exchange ratio; $\dot{VE}$: minute ventilation; OP: oxygen pulse; AT: anaerobic threshold; P values are shown for between groups comparisons (ANCOVA).
Table 3. Relationship between CPET variables and post-operative recovery outcomes.

<table>
<thead>
<tr>
<th></th>
<th>Peak $\dot{V}O_2$</th>
<th>Peak-OP</th>
<th>Peak $\dot{V}E$</th>
<th>PPO</th>
<th>AT</th>
</tr>
</thead>
<tbody>
<tr>
<td>HDU length of stay</td>
<td>-0.562**</td>
<td>-0.372**</td>
<td>-0.421**</td>
<td>-0.514**</td>
<td>-0.454**</td>
</tr>
<tr>
<td>Hospital length of stay</td>
<td>-0.560**</td>
<td>-0.465**</td>
<td>-0.298*</td>
<td>-0.457**</td>
<td>-0.360*</td>
</tr>
</tbody>
</table>

Peak $\dot{V}O_2$: peak rate of oxygen consumption; Peak-OP: peak oxygen pulse; Peak $\dot{V}E$: peak ventilatory volume; PPO: peak power output; $P$ values are shown for Spearman’s Rank Correlation Coefficient: * $P < 0.05$; ** $P < 0.001$