Continuous and intermittent exercise responses in individuals with chronic obstructive pulmonary disease

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Background: While the acute physiological responses to continuous exercise have been well documented in individuals with chronic obstructive pulmonary disease (COPD), no previous study has examined the response to intermittent exercise in these patients.

Methods: We examined the physiological responses of 10 individuals with moderate COPD (forced expiratory volume in 1 second 52 (15)% predicted) who performed both an intermittent (1 min exercise and rest intervals) and a continuous cycle ergometer test on separate days. Both intermittent and continuous exercise tests were performed at the same power output, calculated as 70% of the peak power attained during an incremental exercise test.

Results: Intermittent exercise was associated with significantly lower values for oxygen uptake, carbon dioxide output, expired ventilation, heart rate, plasma lactate concentration, and ratings of breathlessness than continuous exercise. Subjects were able to complete a significantly greater total amount of work during intermittent exercise (71 (32) kJ) than during continuous exercise (31 (24) kJ). The degree of dynamic lung hyperinflation (change in end expiratory lung volume) was significantly lower during intermittent exercise (0.23 (0.07) l) than in continuous exercise (0.52 (0.13) l).

Conclusions: The greater amount of work performed and lower measured physiological responses achieved with intermittent exercise may allow for greater peripheral training adaptations in individuals with more limited lung function. The results suggest that intermittent exercise may be superior to continuous exercise as a mode of training for patients with COPD.

Endurance training has been shown to reduce ventilatory demand during exercise and improve peripheral muscle function in patients with chronic obstructive pulmonary disease (COPD). While these results are encouraging, individuals with moderate to severe COPD are often unable to sustain prolonged periods of exercise at intensities that could maximise peripheral muscle and central cardiovascular adaptations. Consequently, intermittent exercise (IE) has been suggested as an alternative training modality that may be better tolerated in patients with COPD. Intermittent exercise is characterised by repeated short bouts of exercise separated by periods of rest and may be more suited to COPD patients whose daily activities typically require short bursts of exertion interspersed with periods of recovery.

In healthy individuals IE is associated with lower values for heart rate, oxygen uptake (VO₂), expired ventilation (VE), and blood lactate concentration than continuous exercise (CE). An important functional outcome is that total work performed is greatly increased during IE compared with CE when performed at the same absolute intensity. Moreover, Morris and colleagues showed that the lower VO₂ response associated with IE compared with CE performed at 70% of peak power was primarily due to the exponential VO₂ kinetic response typical of a step change in activity level (on-transient VO₂ kinetics). At the onset of constant load exercise a transient exponential increase in VO₂ is observed, attaining a new steady state level within 120–180 seconds during light to moderate intensity exercise. Even during higher intensity exercise that engenders a slow component of increasing VO₂, the on-transient kinetic response remains exponential. Thus, for exercise intervals greater than 120 seconds, the VO₂ attained at the end of each exercise interval approaches that reached during CE at the same exercise intensity. This suggests that the duration of the exercise period is an important factor in determining the IE response, with longer exercise intervals reducing the differences observed between CE and IE.

To date there have been no studies examining the acute response to IE in individuals with COPD. However, Vogiatzis et al reported that the perception of dyspnoea was systematically lower during IE than during CE training in patients with moderate to severe COPD, despite IE being performed at a greater relative intensity than CE. A reduced ventilatory demand and lower degree of dynamic lung hyperinflation during IE could contribute to a reduction in dyspnoea and an improvement in exercise tolerance, independent of the metabolic demand for this mode of exercise. Studies in patients with COPD have also demonstrated slowed on-transient VO₂ kinetics (measured as the time constant or time taken to attain 63% of the asymptotic amplitude) compared with healthy control subjects during submaximal exercise. Slower kinetics may in turn result in a relatively lower VO₂ amplitude during IE.

Severe deconditioning and dyspnoea with exertion are major deterrents to physical activity in COPD patients and could reduce compliance with exercise rehabilitation programmes. If total work performed and/or time to exhaustion are increased and perceived intensity of breathlessness reduced during IE compared with CE, then IE may be a useful training mode in the pulmonary rehabilitation setting. The purpose of the present study was therefore to compare the physiological responses to IE using 1 minute exercise and 1 minute rest intervals, and CE performed at the same intensity.

Abbreviations: CE, continuous exercise; EELV, end expiratory lung volume; FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity; FRC, functional residual capacity; HRpeak, peak exercise heart rate; IC, inspiratory capacity; IE, intermittent exercise; TLC, carbon monoxide transfer factor; TLCO, carbon monoxide transfer factor; V̇Epeak, peak O₂ uptake; V̇O₂peak, peak O₂ uptake; V̇Epeak, peak expired ventilation
absolute power output in a group of COPD patients with moderate airflow obstruction. Performing IE and CE at the same absolute intensity will normalise the metabolic demand and allow for a comparison of the physiological responses between the two exercise modes. It is hypothesised that (1) exercise tolerance (total work performed) will be greater during IE than during CE, (2) IE will be associated with a lower degree of physiological perturbation (V̇O₂, V̇E, blood lactate concentration, dyspnoea), (3) the degree of dynamic hyperinflation will be lower in IE than in CE, and (4) dynamic hyperinflation will be positively correlated with breathlessness.

**METHODS**

**Subjects and experimental design**

Five men and five women with COPD of mean (SD) age 68 (8) years participated in this study. Inclusion criteria for the study were (1) patients classified as having moderate COPD,³⁵ (2) shortness of breath on exertion, and (3) no documented history of substantial co-morbidity. Subjects visited the laboratory on four separate occasions with each visit separated by at least 48 hours. During the first visit the subjects performed pulmonary function tests, were familiarised with the exercise testing procedures, and provided written informed consent. The second visit was used to determine each subject’s maximal exercise capacity. During the subsequent two visits the subjects performed CE and IE on a cycle ergometer. The order of the CE and IE tests was randomised, and the subjects performed only a single bout of exercise on either test day. The experimental protocol was approved by the Griffith University Human Research ethics committee.

**Experimental procedures**

**Pulmonary function assessment**

Spirometry, static lung volumes, lung transfer factor (TlCO), and inspiratory capacity (IC) during exercise were measured using a closed circuit pulmonary function testing system (Collins GS Modular PFT, Braintree, MA, USA). Total lung capacity (TLC) was measured using the helium dilution method while TlCO was assessed with the single breath carbon monoxide test.

**Determination of peak V̇O₂ for cycling**

The incremental exercise test used to measure peak V̇O₂ was performed on a Lode cycle ergometer (Excalibur Sport, Groningen, The Netherlands). Subjects commenced unloaded cycling for 3 minutes and then the power output was increased by 4 W/30 s for men and by 3 W/30 s for women until volitional termination of the test.

During the incremental cycling test subjects breathed through a mouthpiece and wore a nose clip. V̇O₂, carbon dioxide output (V̇CO₂) and V̇E were measured breath by breath while TLC was assessed with the single breath capacity (TLC) was measured using the helium dilution method. Total lung capacity and inspiratory capacity (IC) during exercise were measured using a closed circuit pulmonary function testing system and inspiratory capacity (IC) during exercise were measured during exercise tests. The tests were terminated if subjects were able to complete 30 minutes of CE and/or 60 minutes of IE. Gas exchange indices were measured as described for the incremental exercise test. The breath by breath data were smoothed using a middle five of seven breath averaging procedure. Heart rate and rhythm (Lohmeier M607, Munich, Germany) were monitored during the tests with the ECG electrodes placed in a CM5 configuration. Gas exchange and heart rate values are reported as the peak value attained within a 10 second bin at the end of every minute of exercise. For example, the value at the end of the first minute of exercise for both modes would represent the peak value attained between 0:50 and 1:00 min:s. Total work completed was calculated as the product of exercise time and external power output.

**Effect of V̇O₂ kinetics on intermittent exercise response**

In order to determine if any differences in V̇O₂ between the exercise modes were related to the on-transient kinetics of V̇O₂, the time constant (τ) of the initial rise in V̇O₂ (the phase II or primary component) during the CE bout was calculated (see online data supplement at www.thoraxjnl.com/. supplemental). The phase II τ (τ = 82 (8) seconds, N = 7) was then used to predict the V̇O₂ amplitude at 1 minute of exercise and compared with the mean measured V̇O₂ obtained during the exercise intervals of the IE bout.¹⁰ To demonstrate the effect of the speed of the on-transient component on the V̇O₂ amplitude for IE, a τ value of 42 seconds was also used to predict the IE response. This τ value was previously determined for healthy older subjects performing constant load exercise at a similar relative intensity to that used in the present study.¹⁰

**Determination of plasma lactate concentration**

Before commencement of the CE and IE tests an indwelling cannula was inserted into an antecubital vein of each subject. The plasma lactate concentration was determined (Ciba-Corning Blood Gas Analyser, Medfield, MA, USA) from blood samples obtained at rest, at the end of 3 minutes of unloaded cycling, after 3 and 6 minutes of CE and IE, and at the end of the exercise bouts.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Subject characteristics, pulmonary function, and peak exercise values obtained during incremental exercise</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>68 (8)</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>165 (12)</td>
</tr>
<tr>
<td>Body mass (kg)</td>
<td>68.7 (14.0)</td>
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<tr>
<td>Pulmonary function</td>
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<tr>
<td>FEV₁ (l)</td>
<td>3.15 (0.50)</td>
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<tr>
<td>FVC (l)</td>
<td>2.86 (1.15)</td>
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<tr>
<td>FEV₁/FVC (%)</td>
<td>48 (7)</td>
</tr>
<tr>
<td>TLC (l)</td>
<td>6.64 (1.87)</td>
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<tr>
<td>TlCO (ml/min/mmHg)</td>
<td>9.32 (0.76)</td>
</tr>
<tr>
<td>Heart rate (beats/min)</td>
<td>126 (13)</td>
</tr>
<tr>
<td>Incremental exercise test</td>
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</tr>
<tr>
<td>V̇O₂peak (ml/kg/min)</td>
<td>14.8 (2.3)</td>
</tr>
<tr>
<td>Peak power (W)</td>
<td>57 (25)</td>
</tr>
<tr>
<td>V̇peak (l/min)</td>
<td>38.4 (15.3)</td>
</tr>
<tr>
<td>HR peak (beats/min)</td>
<td>126 (13)</td>
</tr>
</tbody>
</table>

Values presented are mean (SD). FEV₁: forced expiratory volume in 1 second; FVC: forced vital capacity; TLC: total lung capacity; FRC: functional residual capacity; TlCO: carbon monoxide transfer factor; V̇O₂peak: peak O₂ uptake; V̇peak: peak expired ventilation; HRpeak: peak exercise heart rate.
Measurement of inspiratory capacity and end expiratory lung volume

End expiratory lung volume (EELV) was calculated by subtracting IC from the TLC. Inspiratory capacity was measured at rest (while seated on the cycle ergometer), following 7 minutes of exercise, and at the end of CE and IE. The IC manoeuvre was demonstrated before each test by one of the investigators. At each measurement point the subjects were prompted and encouraged to inspire maximally to TLC. The sampling lines to the pulmonary function system and the pneumotachograph and gas sampling lines to the metabolic cart were interfaced through a single mouthpiece worn throughout the duration of the CE and IE tests.

Ratings of breathlessness

During the CE and IE tests subjects were asked to provide ratings of their perceived shortness of breath using a word labelled visual analogue scale. Following a clear explanation of the scale, the subjects were asked to rate their breathlessness at rest, following 7 minutes of exercise, and at the end of exercise.

Statistical analysis

All results are presented as mean (SE) unless otherwise stated. A two way analysis of variance was performed to detect changes in the dependent variables across the two within subject factors (time and exercise mode). Post hoc tests with Bonferroni adjustments were used when a significant interaction or main effect was identified. The differences in time to exhaustion, total work completed, EELV, and subjects’ rating of breathlessness between the two exercise modes were compared using dependent samples t tests. The relationship between EELV and perceived intensity of breathlessness was assessed using Pearson’s correlation coefficient. Statistical significance was accepted at p<0.05. Data were analysed using SPSS Version 11.5 (Chicago, IL, USA).

RESULTS

The physical characteristics of the subjects, pulmonary function test results, and peak exercise values obtained during incremental cycling are shown in table 1. The pulmonary function results showed that the subjects had moderate COPD. Peak VO₂ was reduced by about 40% relative to normal age predicted values.

Figure 1 (A) Oxygen uptake, (B) CO₂ output, (C) expired ventilation and (D) heart rate responses during continuous (closed circles) and intermittent exercise (open circles). Error bars denote SE. The rest intervals were eliminated from the IE trend to equalise the timeline between exercise modes. UL: unloaded cycling; EE: end exercise. *p<0.05, continuous exercise v intermittent exercise.

Figure 2 Plasma lactate response during continuous (closed circles) and intermittent exercise (open circles). Error bars denote SE. UL: unloaded cycling; EE: end exercise. The rest intervals were eliminated from the IE trend to equalise the timeline between exercise modes. *p<0.05, continuous exercise v intermittent exercise.
Intermittent exercise in COPD

Figure 1 shows the cardiopulmonary responses measured during CE and IE. A significant mode by time interaction was observed for VO₂ (F = 41.87, p < 0.001), VCO₂ (F = 55.30, p < 0.001), VE (F = 21.57, p < 0.001) and heart rate (F = 24.43, p < 0.001). For both modes of exercise VO₂, VCO₂ and VE increased significantly from the unloaded to loaded exercise transition (measured 1 minute after the workload was applied). These variables remained relatively unchanged throughout the duration of IE, but continued to increase significantly up to 6 minutes of CE. During CE the end exercise values for VO₂, VCO₂ and VE were not significantly different from the 6 minute values. There was a significant increase in heart rate from unloaded cycling to the end of exercise in both CE and IE (fig 1D). However, the increase in heart rate between consecutive time points was only significant from 3 minutes of CE onwards and between 6 minutes and the end of exercise during IE.

Continuous exercise resulted in significantly higher values for VO₂, VCO₂, VE and heart rate from 3 minutes of exercise onwards in comparison to IE. At the end of exercise the mean differences between CE and IE were: VO₂ 0.25 l/min (95% CI 0.15 to 0.34); VCO₂ 0.34 l/min (95% CI 0.23 to 0.45); VE 9.14 l/min (95% CI 4.94 to 13.30); heart rate 15 beats/min (95% CI 7 to 21). The predicted VO₂ (0.85 (0.09) l/min) using τ values calculated from the individual CE tests was not significantly different from the values measured during IE (0.82 (0.07) l/min). However, experimentally “speeding” on-transient kinetics through the use of a τ value derived from healthy older individuals resulted in predicted VO₂ values (0.97 (0.10) l/min) that were significantly greater than the values measured in our subjects.

A significant mode by time interaction was also observed for plasma lactate concentrations (F = 8.31, p < 0.001). Plasma lactate increased from rest through to the end of exercise during both CE and IE (fig 2). While plasma lactate concentrations during CE were systematically greater than during IE, the differences between exercise modes were only significant at 6 minutes and onwards. The increase in plasma lactate concentrations from rest to the end of exercise was significantly greater during CE (3.1 (0.7) mmol/l) than during IE (1.8 (0.4) mmol/l). The mean difference between exercise modes for plasma lactate concentration at end exercise was 1.7 mmol/l (95% CI 0.8 to 2.5).

The subjects were able to exercise longer and to complete a greater amount of work during IE than during CE. Since the majority of the subjects attained the 60 minute time limit during IE but none achieved the 30 minute CE limit, there was potential for an even greater amount of work to be completed during IE. Intermittent exercise was associated with a lower measured physiological response, with VO₂, VCO₂, VE, heart rate, and plasma lactate concentrations all systematically lower than in CE. Moreover, subjects had a lower level of dynamic hyperinflation during IE than during CE, as assessed by the change in EELV, and this may in turn have contributed to their lower sensation of breathlessness. The results of the present study indicate that IE is better tolerated than CE in patients with COPD because of improved ventilatory mechanics as well as reduced metabolic and cardiovascular perturbations.

As previously discussed, the amplitude of the VO₂ response at a given exercise intensity during IE is determined primarily by the duration of the exercise period. The 1 minute intervals used during IE in the present study would therefore result in lower VO₂ as well as VCO₂, VE and heart rate values, given that these variables are tightly coupled to metabolism within the active muscle during exercise. When exponential models were applied to the VO₂ data measured during the CE bouts, the predicted VO₂ values at the end of 1 minute of exercise were not significantly different from the VO₂ values obtained during the exercise intervals of the IE bouts (see online data supplement at www.thoraxjnl.com-supplemental). Moreover, the measured VO₂ values were not significantly different between exercise modes during the first minute of exercise following the application of the predetermined workload.
that the lower VO2 values observed during IE compared with transition from lower to higher exercise intensities in these anaerobic glycolysis at the onset of exercise or during a result in a greater reliance on phosphocreatine stores and may then be mediated primarily by an increase in breathing demand during exercise results in the progressive trapping of excess non-metabolic CO2, which would be disadvantageous to respiratory health. During IE it has been shown that the lower VO2 values observed during IE compared with CE may also reflect a slower than normal on-transient response in patients with COPD than in healthy subjects. The greater plasma lactate response associated with CE during IE compared with CE suggest either a lower degree of anaerobic metabolism to fuel the energetic demands for IE, is determined by peripheral factors. Of benefit to individuals whose exercise tolerance is limited by peripheral factors.

In summary, we have shown that individuals with COPD, moderate exercise intolerance in patients with COPD.19,20 During IE it has been shown that lactate production occurs primarily during the exercise period21 while the rest period is associated with net lactate removal.22 Thus, despite the possibility of a greater reliance on anaerobic metabolism to fuel the energetic demands for IE, the short exercise intervals interspersed with rest periods resulted in systematically lower plasma lactate values during IE. The greater plasma lactate response associated with CE than with IE may also have contributed, in part, to the higher VE observed during CE. The buffering of hydrogen ions linked with lactic acid production ultimately results in the formation of "non-metabolic" CO2.23 Both an increase in hydrogen ion concentration and excess of "non-metabolic" CO2 production act to stimulate ventilation.24 Since exertional dyspnoea can be a major contributing factor to exercise limitation in patients with COPD, we decided to examine ventilatory mechanics during CE and IE. In many individuals with COPD the increasing ventilatory demand during exercise results in the progressive trapping of air due to incomplete lung emptying.2 The accompanying increase in EELV is termed dynamic hyperinflation. This increase in EELV during exercise, compounded by an already increased resting functional residual capacity (table 1), curtails the ability of the COPD patient to expand tidal volume in line with the greater ventilatory demand.25 An increase in VE must then be mediated primarily by an increase in breathing frequency, which reduces expiratory flow time and further exacerbates the degree of dynamic hyperinflation. Dynamic hyperinflation has been associated with an increase in dyspnoea and may play an important role in the observed exercise intolerance in patients with COPD.25,26 In the present study the degree of dynamic hyperinflation was significantly lower during IE than in CE. Moreover, EELV showed a significant positive correlation with perceived breathlessness. Given the dependence of dynamic hyperinflation on VE, it seems reasonable to suggest that the lower level of perceived breathlessness and dynamic hyperinflation observed during IE was primarily a function of the lower VE during this mode of exercise. In addition, the rest intervals may have provided the subjects with the opportunity to "deflate" their lungs, allowing the lung volume to return to or approach passive FRC before the start of the next exercise bout. This would reduce the degree of dynamic hyperinflation achieved in the subsequent exercise bout.

The findings of the present study suggest that further examination of training intensity and volume, as well as exercise mode, could improve exercise prescription practices in pulmonary rehabilitation programmes. In particular, the use of IE training at intensities comparable to those used in CE protocols may result in similar levels of functional improvement. Recently, Morris et al27 and Vogiatzis et al28 compared CE and IE training in healthy older individuals and in patients with COPD, respectively. These studies showed that, when total work performed was equalised between the exercise modes, the benefits gained were independent of exercise mode as well as intensity. Furthermore, it is our opinion that IE training could possibly benefit individuals ranging in disease severity. For those individuals with severe airflow limitation, breathlessness can be a major psychological deterrent to the pursuit of an active lifestyle.24 The decreased perception of breathlessness associated with IE may improve adherence to training programmes and encourage individuals with lung disease to reverse the debilitating effects of a sedentary lifestyle. The greater options associated with IE protocol design (exercise to rest ratios as well as a greater range of tolerated intensities) could benefit those with mild disease. Even if lung mechanics were not a limiting factor, there is evidence that metabolic and morphological changes in the skeletal muscles of patients with COPD may result in a reduced exercise tolerance.2 In the present study the lower measured levels of plasma lactate during IE compared with CE suggest either a lower degree of metabolic perturbation during IE or greater lactate clearance (during the rest intervals) with better maintenance of intramuscular as well as blood homeostasis. This would be of benefit to individuals whose exercise tolerance is limited by peripheral factors.

In summary, we have shown that individuals with moderate COPD were able to complete a greater amount of work during IE than during CE performed at 70% of peak power. Measures of gas exchange, plasma lactate, dynamic hyperinflation, and perceived breathlessness were significantly lower during IE than during CE. The greater amount of work performed and lower measured physiological responses achieved with IE may allow for greater peripheral training adaptations in individuals with more limited lung function. The results suggest that IE may be superior to CE as a mode of training for patients with COPD.
Adjunct prednisolone therapy does not improve survival in HIV-associated tuberculous pleurisy

Glucocorticoids reduce excessive inflammation in pleural tuberculosis and have been shown to hasten resolution in HIV negative patients. Active tuberculosis can speed progression of HIV infection since HIV replicates more rapidly in activated lymphocytes. Given that prednisolone therapy is associated with reduced immune activation in HIV positive subjects, the authors hypothesised that adjunct therapy with prednisolone in HIV positive patients with pleural tuberculosis would decrease viral replication and improve survival.

In this study, 197 Ugandan HIV infected patients with pleural tuberculosis were randomised to receive treatment for 8 weeks with either a reducing dose of prednisolone or placebo, together with 6 months of treatment with a four-drug antituberculous regime. The median follow up periods were 1.65 and 1.48 years in the prednisolone and placebo groups, respectively. Prednisolone had no effect on either survival or HIV viral load. The mortality rate ratio for prednisolone compared with placebo, after adjusting for confounding factors, was 0.99 (95% CI 0.62 to 1.56, p = 0.95). Use of prednisolone was associated with significantly more rapid improvements in anorexia, weight loss, cough, and radiological resolution of pleural effusion. However, use of prednisolone was also associated with a significantly increased risk of adverse effects (9%) compared with placebo (2%; p = 0.03), and a significantly higher incidence of Kaposi’s sarcoma (prednisolone 4.2 cases/100 person-years; placebo 0 cases/100 person-years, p = 0.02).

In light of the lack of survival benefit and the increased risk of Kaposi’s sarcoma, the authors recommend that prednisolone should not be used in HIV associated tuberculous pleurisy.

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