Assessment of fitness in patients with cystic fibrosis and mild lung disease

P McLoughlin, D McKeogh, P Byrne, G Finlay, J Hayes, M X FitzGerald

Abstract

Background – Maximal exercise testing is used in patients with cystic fibrosis to assess functional status and prognosis. The lactate threshold is an index of aerobic fitness with significant advantages over maximal exercise tests. This study was undertaken to determine if the lactate threshold might be identified, non-invasively, in adult patients with cystic fibrosis and mild lung disease by measurement of ventilatory and gas exchange parameters.

Methods – Ten subjects with mild cystic fibrosis (forced vital capacity (FVC) >70% predicted) and 10 healthy controls undertook an incremental exercise test on a bicycle ergometer. Ventilation and gas exchange parameters were measured continually and arterialized venous blood pH, carbon dioxide tension (PCO₂), and lactate concentrations were measured at intervals throughout the tests.

Results – In subjects with cystic fibrosis there was no significant difference between the mean gas exchange and lactate thresholds (mean difference 1.0 (95% confidence interval (CI) of the mean −1.5 to 3.44) ml/kg/min). In contrast, there was a significant difference between the mean ventilatory and lactate thresholds (3.8 (95% CI 0.9 to 6.7) ml/kg/min). Arterialized venous PCO₂ increased significantly during the exercise tests. In healthy subjects the mean differences between these thresholds were not significantly different from zero and PCO₂ fell significantly during the tests.

Conclusions – The ventilatory threshold significantly overestimates the lactate threshold in subjects with cystic fibrosis induced lung disease because of impaired carbon dioxide excretion during exercise. However, the gas exchange threshold may be used to determine the lactate threshold in this patient group.

(Keywords: cystic fibrosis, exercise, lactate threshold, gas exchange threshold)

Regular aerobic exercise can form an important part of the treatment programme of patients with cystic fibrosis.¹ Such exercise promotes sputum clearance,² ⁴ improves functional status,⁵ ⁶ and fosters a feeling of well-being.⁷ More recently it has been reported that prolonged survival in these patients is associated with higher levels of performance in formal exercise tests, leading to the suggestion that regular aerobic training may increase life expectancy.⁸ To investigate this important issue, improved methods of quantifying aerobic fitness are required which are not excessively stressful and are readily acceptable to patients.

Various techniques have been used to assess fitness in subjects with cystic fibrosis including maximal oxygen consumption, maximum power output, time to exhaustion during incremental exercise tests, time to exhaustion at a fixed work load, or time taken to complete a fixed distance.⁵⁶⁷ ⁹ However, there are many difficulties associated with their use in this context.¹⁰ In particular, all require the subject to make maximal efforts, rendering such tests highly dependent on motivation.

The lactate threshold is an alternative index of aerobic fitness which has been widely used in athletes. As the workload progressively increases during an incremental exercise test, a critical exercise intensity is reached above which blood lactate levels increase progressively, reflecting an abrupt rise in lactate concentration in the exercising muscles.¹⁰ The oxygen uptake of the subject at this critical work load is called the lactate threshold (LT), which is a better indicator of aerobic fitness than other indices commonly used. It permits identification of the optimum training intensity for an individual and does not require exercise to exhaustion for its determination.¹⁰ ¹⁴ In healthy subjects the LT may be determined non-invasively by identification of the ventilatory threshold (Ve T) which is defined as the point during an incremental exercise test when ventilation breaks away from its initial linear relationship to oxygen uptake (VO₂).¹⁰ Subjects with significant obstructive lung disease frequently are unable to produce this hyperventilatory response. However, in these patients it is possible to detect a non-linear increase in carbon dioxide output in relation to VO₂ – the gas exchange threshold (GET) – which coincides with the LT.¹³ ¹⁶

The lung disease associated with cystic fibrosis leads to a characteristic pattern of pathological disturbances even at an early stage when spirometric abnormalities are minimal or mild. In particular, there are disproportionately large increases in physiological dead space and shunt.¹⁷ ¹⁸ By impairing carbon dioxide excretion these disturbances might prevent the identification of the ventilatory and gas exchange thresholds.

The primary purpose of the present study was to determine if the LT might be identified non-invasively in adult cystic fibrosis patients with mild to moderate lung disease by measure-
ment of ventilatory and gas exchange parameters and thus provide an easily determined index of aerobic fitness.

Methods

Ten male subjects (>14 years of age) with cystic fibrosis (sweat test demonstrated sodium and chloride concentrations >60 mmol/l) and mild to moderate lung disease (forced vital capacity (FVC) >70% of predicted) were recruited from the Adult Cystic Fibrosis Clinic at St Vincent’s Hospital. Ten non-smoking control subjects who had no history of significant cardio-pulmonary disease were also recruited. The study was approved by the hospital ethics committee and all participants gave written informed consent.

Each subject attended the exercise laboratory on a single occasion and completed an incremental exercise test to exhaustion on a friction braked cycle ergometer (Monark, Sweden). Subjects were instructed to maintain a constant pedalling rate. To facilitate this, their actual pedalling rate was displayed continuously to the subject. The test began with a four minute period of unloaded cycling after which the workload was increased every two minutes by 10–20 W, depending on the stature and exercise history of the subject. They did not come to the laboratory for prior training or acclimatisation.

Subjects wore a nose clip and mouthpiece connected to a non-rebreathing valve. Minute ventilation (V̇e), oxygen uptake (VO₂), and carbon dioxide output (VCO₂) were determined at 15 second intervals using a computerised system (PK Morgan, Kent, UK). Inspiratory flow rates were measured with a turbine flow meter and electrically integrated to give volume. Expired gas passed through a mixing chamber and was then continuously sampled for determination of mixed expired gas composition. Oxygen saturation (Biox 3700E, Ohmeda, Colorado, USA) and an ECG were monitored throughout the exercise tests. Calibrations were checked before and after each test using calibration syringes and precision oxygen and carbon dioxide gas mixes (BOC, Ireland).

Forced expiratory volume in one second (FEV₁) and FVC were measured with a model S spirometer (Vitalograph Ltd, Bucks, UK) before exercise and peak flow was determined both before and after the exercise test. Predicted values of these variables were calculated using the equations of Quanjer et al.¹ Predicted maximum voluntary ventilation (MVV) was calculated as FEV₁ × 37.5.²⁰ Predicted maximum heart rate was calculated as 220 – age in years.²¹

Arterialised venous blood samples were drawn over the last 15 seconds of each workload as previously described.²² A portion of each sample was stored anaerobically on ice for subsequent measurement of pH and PaCO₂ using an automated self-calibrating blood gas analyser (Model IL 1312 Blood Gas Manager; Allied Instruments Lab, Cheshire, UK). 1.0 ml of the remaining blood was pipetted into 2.0 ml of chilled 7% perchloric acid and the supernatant assayed for lactate concentration using an enzymatic, spectrophotometric technique (Sigma Diagnostics Procedure No. 826-uv; Sigma Laboratories, Dorset, UK).

Calculations

All determinations of lactate, ventilatory, and gas exchange thresholds were performed by a blinded observer. The GET was identified using the modified V-slope method of Sue et al.²³ and the VenT was determined by identifying the point at which the ventilatory equivalents for oxygen began a steady progressive rise after a period when they had been unchanging or progressively falling.²⁴ The LT was determined using the method of Beaver et al.²⁵ All thresholds were expressed as oxygen uptake (ml/kg/min).

Figure 1 illustrates the method of determining the GET in a single subject. VCO₂ is plotted against VO₂ and a line parallel to the line of identity is plotted through the lower data points. This is most conveniently accomplished by drawing the x and y axes of the VCO₂ versus VO₂ plot to an exactly equal scale, ensuring that the line of identity makes an angle of 45° (slope of 1) with the x axis. A 45° set square is then placed on the x axis so that so that its angled edge lies parallel to the line of identity. By moving the set square horizontally along the x axis a best-fit line may be drawn through the lower data points by eye. This manoeuvre identifies the point during the exercise test, expressed as VO₂ (ml/kg/min), when VCO₂ begins to rise disproportionately faster than VO₂ due to a sudden increase in lactic acid concentration within the exercising muscles. The slope of a line through the data points below the lactate threshold is approximately equal to 1 because exercising muscle primarily uses glycogen as a source of energy. As a consequence, the ratio of the increase in VCO₂ to the increase in VO₂ is approximately 1.²⁴
Lactate threshold in cystic fibrosis

Figure 2 Determination of the lactate threshold (LT) for the same subject as in fig 1. The logarithm of blood lactate concentration was plotted against the logarithm of the mean VO$_2$ during the minute in which that blood sample was drawn. The lactate threshold was determined as the point of intersection of the two least squares regression lines fitted to the data points before and after the onset of a rapid increase in lactate concentration (chosen by blinded reviewer). See text for details.

Figure 2 demonstrates the method of identification of the LT using data obtained from the same subject as in fig 1. Log lactate concentration is plotted against log VO$_2$ and the point of inflection was chosen by a blinded reviewer. Using least squares regression a straight line was fitted to the data points below and including the chosen inflection point and a separate line to the data points above and including the inflection point. The intersection of these two lines indicates the LT which is expressed as VO$_2$ (ml/kg/min).$^{23}$

**Table 1 Mean (SD) anthropometric characteristics, pulmonary function and measured blood parameters at rest in patient and control groups (n = 10)**

<table>
<thead>
<tr>
<th></th>
<th>Patients</th>
<th>Controls</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>23.4 (4.3)</td>
<td>24.1 (4.4)</td>
</tr>
<tr>
<td>Height (m)</td>
<td>1.75 (0.05)*</td>
<td>1.80 (0.05)</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>65.8 (9.0)**</td>
<td>80.4 (10.8)</td>
</tr>
<tr>
<td>FEV$_1$, (% predicted)</td>
<td>80.4 (15.6)**</td>
<td>103.9 (11.0)</td>
</tr>
<tr>
<td>FVC, (% predicted)</td>
<td>96.9 (15.8)</td>
<td>107.7 (14.9)</td>
</tr>
<tr>
<td>PEF, (% predicted)</td>
<td>70.4 (9.7)**</td>
<td>81.8 (7.9)</td>
</tr>
<tr>
<td>pH</td>
<td>7.415 (0.018)</td>
<td>7.406 (0.014)</td>
</tr>
<tr>
<td>Pco$_2$, (kPa)</td>
<td>4.9 (0.3)</td>
<td>5.2 (0.3)</td>
</tr>
<tr>
<td>Actual bicarbonate (mmol/l)</td>
<td>23.1 (1.2)</td>
<td>23.9 (1.2)</td>
</tr>
<tr>
<td>Blood lactate (mmol/l)</td>
<td>1.2 (0.3)</td>
<td>0.8 (0.2)</td>
</tr>
</tbody>
</table>

FEV$_1$ = forced expiratory volume in one second; FVC = forced vital capacity; PCO$_2$ = carbon dioxide tension.
*p<0.05, **p<0.01 (unpaired t test) compared with control value.

**Table 2 Mean (SD) blood and ventilatory parameters, heart rate and measured blood parameters at peak exercise in patient and control groups (n = 10)**

<table>
<thead>
<tr>
<th></th>
<th>Patients</th>
<th>Controls</th>
</tr>
</thead>
<tbody>
<tr>
<td>Peak VO$_2$ (ml/kg/min)</td>
<td>32.4 (5.7)**</td>
<td>42.5 (6.1)</td>
</tr>
<tr>
<td>Ventilation (% MVV)</td>
<td>63.5 (15.3)</td>
<td>64.7 (9.6)</td>
</tr>
<tr>
<td>Heart rate (% predicted maximum)</td>
<td>94.4 (5.1)</td>
<td>95.6 (8.3)</td>
</tr>
<tr>
<td>pH</td>
<td>7.290 (0.023)</td>
<td>7.272 (0.042)</td>
</tr>
<tr>
<td>PCO$_2$, (kPa)</td>
<td>5.3 (0.9)</td>
<td>4.8 (0.9)</td>
</tr>
<tr>
<td>Actual bicarbonate (mmol/l)</td>
<td>16.6 (1.7)**</td>
<td>16.1 (1.7)</td>
</tr>
<tr>
<td>Blood lactate (mmol/l)</td>
<td>5.4 (1.6)**</td>
<td>8.3 (2.4)</td>
</tr>
</tbody>
</table>

VO$_2$ = oxygen uptakes; MVV = maximum voluntary ventilation; PCO$_2$ = carbon dioxide tension.
*p<0.01 (unpaired t test) compared with control value.

**DATA ANALYSIS**

One group t tests, unpaired t tests or analysis of variance (ANOVA) with repeated measures were used as appropriate. When the overall ANOVA was statistically significant, Student Neuman Keul’s post hoc test (SNK) was used to assess the significances of the differences between specific means. The method of Bland and Altman was employed to compare the values determined for the anaerobic threshold using the lactate, gas exchange, and ventilatory threshold methods.$^{24}$

**Results**

Anthropometric, pre-exercise pulmonary function, and resting arterialised venous blood data for both patient and control groups are given in table 1. The patients were of lower mean height (p<0.05, unpaired t test) and weight (p<0.01, unpaired t test) than the control subjects and had evidence of obstructive airways disease. The mean (range) Shwachman score for the patient group was 75 (66–85). Table 2 shows values recorded at peak work load. Both patients and controls had developed a marked metabolic acidosis at the end of the exercise test and the blood lactate levels were increased in all participants by more than 3 mmol/l. The mean maximum ventilation at peak exercise in both the cystic fibrosis and control groups was markedly less than the MVV (table 2), while heart rate was near maximal. The peak flow after exercise was not reduced below the pre-exercise values in any of the study participants.

In subjects with cystic fibrosis the mean difference (95% confidence interval of the mean) between the GET and the LT was 1.0 (−1.5 to 3.44) ml/kg/min, not significantly different from zero (p>0.05, one sample t test). This indicates that the GET could be used to provide an unbiased estimate of the LT in a group of patients with cystic fibrosis. The mean difference between the LT and the VenT was 3.8 (0.9 to 6.7) ml/kg/min, a value significantly greater than zero (p<0.05, one sample t test). This suggests that use of the VenT would lead to an overestimate of the LT in a population of cystic fibrosis patients. The corresponding comparisons for the control groups demonstrated a mean difference between LT and GET of −0.1 (−2.5 to 2.3) ml/kg/min and a mean difference between LT and VenT of 1.9 (−1.8 to 5.5) ml/kg/min, neither of which are significantly different from zero (p>0.05, one sample t test).

In order to illustrate the agreement of the GET and LT within individual subjects, a comparison of the two thresholds in the patient group is shown in fig 3 using the technique of Bland and Altman.$^{24}$ For each subject the difference between the LT and GET is plotted against the average of the two determinations. The limits of agreement (mean ± 2×SD of differences) are also shown (−6.8 to 8.8 ml/ kg/min). For the control group the limits of agreement were similar (−7.6 to 7.6).

We examined the behaviour of arterialised venous PCO$_2$ throughout the incremental tests (fig 4). For each subject the value of arterialised
groups was then plotted against the cor-
may contribute to hypercapnia is the presence
work load completed (Peak WL). The mean frequency of breathing to ventilatory stimuli
half way between the LT and the maximum
patients with cystic ®brosis abnormalities in
work load and the LT (0.5 LT), (c) at the LT,
d) at a work load which elicited a V\text{O}_2 \text{ that lay}
half way between the LT and the maximum
work load (LT+0.5\text{A}), and (e) at the peak
work load completed (Peak WL). The mean
value of these variables for patient and control
groups was then plotted against the corre-
spending ®ndings may have arisen because
previous workers used end tidal carbon dioxide
as an index of arterial carbon dioxide whereas
in the present study we have measured blood
arterialised venous P\text{CO}_2.

Discussion

The subjects with cystic fibrosis did not
hypoventilate normally following the onset
of lactic acidosis, as shown by the absence of a
fall in arterialised venous P\text{CO}_2 at heavier work
loads (fig 4). This ®nding is in contrast to
previous reports that patients with mild to mod-
erate cystic fibrosis induced lung disease do
not retain carbon dioxide during exercise.\textsuperscript{25,26}
The differing ®ndings may have arisen because
previous workers used end tidal carbon dioxide
as an index of arterial carbon dioxide whereas
in the present study we have measured blood
arterialised venous P\text{CO}_2 directly.\textsuperscript{25,26}

The impaired carbon dioxide excretion
which we observed was not due to ventilatory
limitation since the mean (SD) maximum min-
ute ventilation of these patients was only 63.5
(15.3) % of the predicted maximum voluntary
ventilation, whereas the mean maximum heart
rates were greater than 90% predicted max-
imum in both groups (table 2). A mechanism
which may have contributed to the impaired
excretion of carbon dioxide during exercise is
an increase in physiological dead space vent-
ilation which has been reported in patients with
cystic fibrosis, even when standard spirometric
tests are only mildly abnormal.\textsuperscript{14} This abnormal
increase in physiological dead space ventil-
ation may arise because of abnormalities of ven-
tilation-perfusion matching in the lung. In
patients with cystic fibrosis abnormalities in
the pattern of response of tidal volume and
frequency of breathing to ventilatory stimuli
may also contribute.\textsuperscript{27} A second factor which
may contribute to hypercapnia is the presence
of obstructive airways disease (table 1). Sub-

venous P\text{CO}_2 and lactate was determined
during (a) unloaded cycling (“0” WL), (b) at a work
load which elicited a V\text{O}_2 half way between “0”
work load and the LT (0.5 LT), (c) at the LT,
(d) at a work load which elicited a V\text{O}_2 that lay
half way between the LT and the maximum
work load (LT+0.5\text{A}), and (e) at the peak
work load completed (Peak WL). The mean
value of these variables for patient and control
groups was then plotted against the corre-
spending line, is not signi®cantly
different from zero (p>0.05, unpaired t test).
Upper and lower lines indicate the limits of agreement (mean ±
2.262 × SD of the differences).
Lactate threshold in cystic fibrosis

projects with well controlled asthma and healthy subjects breathing through an added resistance fail to show normal hyperventilation at heavier work loads. The impaired carbon dioxide excretion at higher work loads is not due to a blunted central ventilatory response to hypercapnia since this is normal in patients with mild to moderate cystic fibrosis induced lung disease. All of the subjects with cystic fibrosis were able to continue to a work load which caused a significant metabolic acidosis (blood lactate >3 mmol/l). In addition, none of them demonstrated a lactate threshold until more than two minutes of the incremental exercise test had elapsed, so the gas exchange threshold occurred at a time when carbon dioxide and oxygen stores were changing at a constant rate. These are the necessary conditions which must be met to allow identification of the gas exchange threshold. In patients with more severe lung disease these criteria may not be satisfied. Thus, to determine if the findings of the present study can be extended to those in whom FVC is less than 70% predicted, further work is required.

We have found that, in patients with cystic fibrosis and mild lung disease, the gas exchange threshold provides an unbiased estimate of the lactate threshold. The standard deviation of the differences in the patient group (3.5 ml/kg/min) was identical to that of the controls (3.5 ml/kg/min), indicating that the technique is equally precise in both patients with mild cystic fibrosis and healthy subjects. Furthermore, the standard deviation is similar to those of previously published studies of patients with smoking related lung disease. To compare our data with previous reports we converted the oxygen consumptions at the gas exchange thresholds to l/min. The resultant standard deviation of the differences (0.230 l/min) was identical to that reported by Dickstein and colleagues in subjects with chronic obstructive pulmonary disease. In another study of subjects with chronic obstructive pulmonary disease Sue et al compared gas exchange and bicarbonate thresholds and observed a value of 0.215 l/min for the standard deviation of the differences, a value similar to that of the present study.

The reproducibility of the determination of the gas exchange and lactate thresholds is clearly relevant to the method comparison which we report here. The reproducibilities of the two methods limit the agreement which is possible between them. To determine the reproducibility of each method would require two tests on each subject separated by a short interval (7–10 days) during which the subjects remained stable. We are unaware of any studies in patients with cystic fibrosis or in normal subjects which examine this issue based on the methods which we have used. Two groups have examined this issue in normal subjects using somewhat different methodologies. Smith and O'Donnell reported a study of reproducibility using a cumulative sum method to determine the ventilatory threshold and found that the mean difference between replicate tests was 0.001 l/min with a standard deviation of 0.220 l/min. Caiozzo et al showed good linear correlation between the values of VO2 at the gas exchange threshold obtained in individual subjects on two separate occasions and found that the standard error of the estimate was 0.260 l/min. Neither group examined the reproducibility of determinations of the lactate threshold. It is interesting to note that the standard deviation of the difference reported by Smith and O'Donnell and the standard error of the estimate reported by Caiozzo et al are similar to the standard deviation of the differences between the gas exchange and lactate thresholds in the present study.

Exercise tests were continued to exhaustion in our study. This protocol was adopted to allow exploration of the ventilatory responses throughout an incremental exercise test. If the sole object of an exercise test is to determine the gas exchange threshold in order to assess aerobic performance in patients with cystic fibrosis offers several further advantages over the commonly employed maximal tests. It is an excellent index of aerobic performance, it can increase following training even in the absence of changes in VO2max, and the changes observed are closely correlated with improved endurance performance. This may be especially important when maximal performance is limited by ventilatory capacity. In this circumstance, training can lead to important cardiac and skeletal muscle effects which lead to improved performance at submaximal work loads. The gas exchange threshold provides an index of performance at such submaximal work loads, information which is directly relevant to a patient’s ability to carry out the tasks of daily living. Finally, the gas exchange threshold allows determination of the optimum training intensity in an individual.

In conclusion, the ventilatory threshold significantly overestimates the lactate threshold in subjects with mild cystic fibrosis induced lung disease because of impaired carbon dioxide excretion during exercise. However, the gas exchange threshold may be used to estimate the lactate threshold in this patient group, both accurately and non-invasively. It provides an index of aerobic fitness which offers significant advantages over previously used methods which depend upon maximal exercise effort.

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