Prolonged Oxygen Kinetics During Constant Workload Submaximal Exercise Is Associated With Disease Severity in Adult Subjects With Cystic Fibrosis

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BACKGROUND: The goal of this study was to explore the relation between oxygen kinetics during constant work load submaximal cardiopulmonary exercise test (CPET) and disease severity in adult subjects with cystic fibrosis. METHODS: Fourteen adult subjects with cystic fibrosis (CF; 8 males, 22 ± 4 y old) and a mean Schwachman score of 73 ± 11 and 10 healthy individuals (5 males, 29 ± 4 y old) underwent pulmonary function tests at rest, maximal and constant work load submaximal CPET on a cycloergometer. Breath-by-breath analysis was used for measuring oxygen kinetic parameters and the time constant (tau), expressing phase 2 of submaximal CPET. RESULTS: Subjects with CF had a significantly prolonged tau compared with healthy subjects (42.3 ± 21.5 vs 29.3 ± 6.4, s, P < .05). The tau during phase 2 was inversely correlated with FEV1(% pred) (r = -0.77, P = .001), breathing reserve (r = -0.74, P = .003), V̇O₂peak (r = -0.53, P = .049), V̇O₂/t slope (r = -0.58, P = .03), and Schwachman score (r = -0.80, P = .001). In a multivariate regression model including all the above variables, the Schwachman score (β = -0.697, P = .002) emerged as independent predictor of tau (R² = 0.719, P = .001). CONCLUSIONS: We conclude that adult subjects with CF present significant prolonged oxygen kinetics during constant work load submaximal exercise in relation to disease severity. Thus, submaximal exercise should be considered the preferable CPET choice in adult patients with severe CF. Key words: cystic fibrosis; submaximal exercise; time constant (tau).

Introduction

Cystic fibrosis (CF) is a progressive, life-limiting disease, with an incidence of 1:2,000 to 1:3,500 among white populations.1 It is caused by mutations in a single gene on the long arm of chromosome 7, which encodes a protein, the cystic fibrosis transmembrane conductance regulator (CFTR).2 More than 1,500 mutations have been identified to be responsible for CFTR quantity or quality defects leading to abnormal chloride concentration across the apical membrane of epithelial cells, especially in the airways and pancreas. This results in thick secretions and, through a vicious cycle of inflammation and infection, in progressive lung disease and malnutrition.2 The disease most commonly affects the respiratory system among the adult population, increasing morbidity and determining the patients’ survival.3,4 The prognosis of the disease is worse in women, and it is dependent on nutritional status, respiratory involvement, colonization by Pseudomonas aeruginosa, and existence of complications such as pneumothorax and right ventricular failure.5 Advances in the diagnosis

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The authors have disclosed no conflicts of interest.

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and the therapeutic management of CF have gradually led to a significant increase in life expectancy, accompanied by a corresponding increase in the survival rate of CF patients.6

Lung function parameters such as FEV₁,7 peak oxygen uptake (VO₂peak) during incremental cardiopulmonary exercise testing (CPET)8 and the clinical evaluation system known as the Schwachman score9 are common prognostic indices used to assess functional capacity and disease severity in CF patients. Peak exercise capacity is reduced in CF,10 and the main responsible factors are diminished pulmonary and nutritional status,11 as well as peripheral muscle weakness.12 Although maximal CPET is a useful clinical tool to measure aerobic capacity directly, it remains difficult for debilitated patients to perform this test, especially CF patients. An alternative modality for estimating aerobic capacity could be oxygen kinetics during submaximal constant work load exercise testing. However, there are very limited data concerning constant work load oxygen kinetics in subjects with CF.13 These data consist mainly of lung function decline and oxygen desaturation during 6-min walk test in children and patients who have received transplants. Furthermore, oxygen kinetics during exercise has been correlated with disease severity only during maximal CPET.

In this study, we hypothesized that (1) adult subjects with CF present impaired oxygen kinetics during constant work load submaximal CPET, and (2) oxygen kinetics is related to disease severity in CF. The aim of this study was to explore oxygen kinetics during constant work load submaximal CPET in adult subjects with cystic fibrosis in comparison with healthy subjects and in relation to disease severity.

Methods

This study involved 14 adult subjects with CF (8 males, mean age: 22 ± 4 y old) referred to the CF out-patient clinic of our Institute and 10 healthy individuals (controls) (5 males, mean age: 29 ± 4 y old). The diagnosis of CF was based on clinical evaluation and laboratory testing, including sweat testing and genotype analysis. All participants were physically active, nonsmokers, and not engaged in regular training. All subjects were clinically stable and optimally treated at the time of the study. Subjects had no clinical symptomatology or signs of cardiac disease, and there was no history of any cardiovascular risk factor for ischemic heart disease. All subjects with CF before referral to our CF out-patient clinic had a routine echocardiogram, which had shown cardiac function within normal limits. Patients and healthy subjects with history of recently (within the previous month) diagnosed pulmonary infection or respiratory failure, cardiac, orthopedic, or neurologic problems affecting exercise capacity were excluded from the study. None of the controls suffered from any acute or chronic disease at the time of testing, or were under medication. The institutional review board approved the study, informed consent was obtained from all subjects, and the study was conducted in accordance with applicable laws and regulations, including the International Conference on Harmonization Guideline for Good Clinical Practice (http://www.ich.org. Accessed January 26, 2015.).

Design of the Study

All subjects with CF and healthy subjects underwent a spirometric evaluation, a symptom-limited CPET, and a constant work load CPET. Subjects with CF were also clinically assessed, and the Schwachman score was calculated.

The latter is a scoring system based on 4 separate aspects of the disease profile: (1) general activity, (2) physical findings (degree of clubbing, auscultatory findings, cough, and sputum), (3) nutritional status, and (4) chest x-ray findings. Each item is given equal weight, from 0 to 25 points. A total of 86 points represents a perfect score,9 whereas a total below 40 points represents a severe condition. The Schwachman score is calculated by 2 different observers to obtain objective results.

Pulmonary Function Assessment

Study participants (healthy and CF subjects) underwent measurement of FVC and FEV₁ in the sitting position with
a closed-circuit spirometer (Sensormedics, Yorba Linda, California) as recommended by the American Thoracic Society. In addition, participants underwent inspiratory capacity (IC) measurement in the sitting position, before exercise, according to previous studies. 

**CPET**

Subjects underwent a symptom-limited, incremental CPET on a cycloergometer (model 2000, Marquette Electronics, Milwaukee, Wisconsin) in a sitting position. Cardiopulmonary data were recorded for 2 min at rest followed by 3 min of unloaded pedaling. The work rate increment was estimated by using the equation of Hansen et al to attain test duration of 8–12 min. A 12-lead electrocardiogram was recorded every minute using the Max 1 system (Marquette Electronics). Blood pressure measurements were made with a standard mercury sphygmomanometer and IC maneuvers were performed every 2 min during exercise and at peak exercise. The IC maneuvers were first explained and then practiced by the subjects until reproducible efforts were made, as previously described and IC peak was measured at the end of exercise. We also measured breathing reserve at maximal exercise as: (MVV – peak VE)/MVV, where MVV was the maximal voluntary ventilation obtained from 40 × FEV$_1$. A pulse oximeter was used for $S_{\text{PO}_2}$ monitoring. The subjects breathed through a mouthpiece with a nose clip in place. Oxygen uptake ($\dot{V}_{\text{O}_2}$), carbon dioxide output ($\dot{V}_{\text{CO}_2}$), and air flow were measured on a breath-by-breath basis using the Vmax 229 monitor for pulmonary and metabolic studies (Sensormedics). The system was calibrated with a gas mixture of known concentration before each test. Measurements were obtained in the sitting position before and during exercise. Baseline $\dot{V}_{\text{O}_2}$ was calculated by averaging the measurements made for 2 min before the beginning of exercise. $\dot{V}_{\text{O}_2}$ was calculated as the average of measurements made during the last 20 s of incremental exercise testing. Anaerobic threshold (AT) was determined using the $V_{\text{slope}}$ technique, and the result confirmed by a graph on which the respiratory equivalent for oxygen ($V_{\text{E}}/V_{\text{O}_2}$) and carbon dioxide ($V_{\text{E}}/V_{\text{CO}_2}$) were plotted simultaneously against time. Heart rate and oxygen pulse ($\dot{V}_{\text{O}_2}$ pulse: $V_{\text{O}_2}$/heart rate) at peak exercise were also calculated.

To evaluate the oxygen uptake kinetics during early recovery, the first degree slope of $\dot{V}_{\text{O}_2}$ for the first 1 min of recovery period was calculated by linear regression using an appropriate computerized statistical program, assuming that the fall in $\dot{V}_{\text{O}_2}$ during early recovery is linear. The ventilatory response to exercise was calculated as the slope by linear regression of $V_{\text{E}}$ versus $V_{\text{CO}_2}$ from the beginning of exercise to AT as in previous studies. Peak work rate was defined as the highest work level reached and maintained at a pedaling frequency of no less than 50 rpm for 20 s. Subjects were instructed to exercise until exhaustion. A respiratory exchange ratio $> 1.09$ was considered to be an indication of a nearly maximal effort.

**Constant Work Load Submaximal CPET**

After 1 h of rest, each study participant underwent a constant load exercise test in a cycloergometer. Initially, the test load was set at 80% of the value of the work rate (WR) corresponding to the AT detected in the maximum test, which was performed first (with a minimum value of 30 W). Cardiopulmonary data were recorded for 2 min at rest, followed by 3 min of unloaded pedaling. After unloaded pedaling, subjects performed the constant load test for 7 min at the work load rate described above, followed by unloaded pedaling for 5 min. We used a 7-min work load for a better evaluation of all oxygen kinetics phases. To improve the confidence of the kinetic parameter determination, this test was performed 3 times, and the averaged profile was used for the kinetics analysis. We also chose to limit the constant work rate to the sub-anaerobic threshold range to obviate the confounding influence of the slow component of the kinetics at work rates associated with a sustained increase in lactate levels. The breath-by-breath measurements of each curve were edited only for breaths that were not reflective of the underlying physiological process interpolated to a second-by-second basis, time-aligned, and averaged to produce a single response. Curve fitting was performed with Origin 7.0 graphics software (Microcal Software, Northampton, Massachusetts) using iterative techniques.

$\dot{V}_{\text{O}_2}$ response was described by a 3-phase model. To characterize phase 1 (cardiodynamic phase), we used the period from the start of the exercise to the point when the respiratory exchange ratio started to fall in conjunction with end-tidal $P_{\text{O}_2}$ starting to increase. During phase 2, $\dot{V}_{\text{O}_2}$ continues to increase as its consumption by muscles increases, and it is defined as the exponential period beginning immediately following phase 1, until steady state (phase 3) at moderate WR levels. The exponential curve of phase 2 is described by the equation: $\Delta y(t) = A[1 - e^{-(t-TD)/\tau}]$, which provides estimates of the time constant (tau). The dependent variable $\Delta y(t)$ equals the increase in oxygen uptake at time t, A equals the increase in oxygen uptake from the onset of loaded exercise to steady-state exercise (phase 3), and TD is the time delay from the onset of loaded exercise until the beginning of the exponential rise in oxygen uptake. There is theoretical and experimental evidence to analyze $\dot{V}_{\text{O}_2}$ responses of submaximal CPET by separating phase 1 and
Table 1. Baseline Demographic Characteristics and Pulmonary Function Assessment at Rest in Subjects With Cystic Fibrosis and Controls

<table>
<thead>
<tr>
<th></th>
<th>Subjects (n = 14)</th>
<th>Controls (n = 10)</th>
<th>P</th>
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</thead>
<tbody>
<tr>
<td>Gender (males)</td>
<td>8</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Age (y)</td>
<td>22 ± 4</td>
<td>29 ± 4</td>
<td>.003</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>21.6 ± 3.2</td>
<td>23.1 ± 2.9</td>
<td>.24</td>
</tr>
<tr>
<td>Schwachman score</td>
<td>73 ± 11</td>
<td>NA</td>
<td></td>
</tr>
<tr>
<td>FEV₁ (% pred)</td>
<td>76.7 ± 31.6</td>
<td>105.9 ± 14.9</td>
<td>.007</td>
</tr>
<tr>
<td>FVC (% pred)</td>
<td>89.2 ± 29.7</td>
<td>106.9 ± 14.3</td>
<td>.07</td>
</tr>
<tr>
<td>FEV₁/FVC (%)</td>
<td>72.9 ± 11.6</td>
<td>86.5 ± 5.6</td>
<td>.001</td>
</tr>
<tr>
<td>IC at baseline (L)</td>
<td>2.6 ± 0.8</td>
<td>2.8 ± 0.6</td>
<td>.18</td>
</tr>
</tbody>
</table>

Continuous variables are presented as mean ± SD.
BMI = body mass index
NA = not applicable
% pred = percentage of predicted
IC = inspiratory capacity

Table 2. Maximal and Submaximal Cardiopulmonary Exercise Testing Parameters in Subjects With Cystic Fibrosis and Controls

<table>
<thead>
<tr>
<th></th>
<th>Subjects (n = 14)</th>
<th>Controls (n = 10)</th>
<th>P</th>
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</thead>
<tbody>
<tr>
<td>Maximal CPET parameters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>VO₂peak (L/min)</td>
<td>1.6 ± 0.4</td>
<td>2.1 ± 0.4</td>
<td>.009</td>
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<tr>
<td>VO₂peak (mL/kg/min)</td>
<td>26.4 ± 6.9</td>
<td>35.4 ± 7.7</td>
<td>.007</td>
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<td>AT (mL/kg/min)</td>
<td>15.2 ± 4.7</td>
<td>25.6 ± 4.8</td>
<td>&lt;.001</td>
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<td>VO₂/t slope (L/min²)</td>
<td>0.74 ± 0.28</td>
<td>0.95 ± 0.19</td>
<td>.048</td>
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<tr>
<td>V̇E/V̇CO₂ slope</td>
<td>30 ± 5</td>
<td>26 ± 3</td>
<td>.046</td>
</tr>
<tr>
<td>O₂ pulse (mL/beat)</td>
<td>9.6 ± 2.4</td>
<td>12.5 ± 2.7</td>
<td>.01</td>
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<tr>
<td>HRpeak (beats/min)</td>
<td>171 ± 12</td>
<td>175 ± 25</td>
<td>.15</td>
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<tr>
<td>ICpeak (L)</td>
<td>2.5 ± 0.7</td>
<td>2.7 ± 0.6</td>
<td>.27</td>
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<tr>
<td>WRpeak (watts)</td>
<td>139 ± 35</td>
<td>150 ± 29</td>
<td>.45</td>
</tr>
<tr>
<td>Phase I (t, s)</td>
<td>29.4 ± 10.2</td>
<td>23.5 ± 6.3</td>
<td>.10</td>
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<tr>
<td>Phase II (tau, s)</td>
<td>42.3 ± 21.5</td>
<td>29.3 ± 6.4</td>
<td>.049</td>
</tr>
</tbody>
</table>

Continuous variables are presented as mean ± SD.
CPET = cardiopulmonary exercise testing parameter
VO₂peak = peak oxygen uptake
AT = anaerobic threshold
VO₂/t slope = first-degree slope of VO₂ for the first 1 min of recovery
V̇E/V̇CO₂ slope = slope of the ventilatory equivalent for carbon dioxide output
HRpeak = heart rate at peak of maximal exercise
ICpeak = inspiratory capacity at peak of maximal exercise
BR = breathing reserve
WRpeak = peak work rate
t = mean duration of phase 1 of submaximal CPET
tau = time constant of phase II during submaximal CPET

phase 2,20,21,24 as phase 1 is the cardiodynamic phase and expresses the abrupt increase in oxygen consumption, whereas the phase 2 expresses mainly the oxygen consumption by muscles.

Statistical Analysis

All continuous variables are presented as mean ± SD. All variables were tested for normal distribution. The unpaired Student t test was used to compare group means of continuous variables of subjects with CF and healthy subjects. The Pearson correlation coefficient was used to test correlations. A univariate linear regression was performed to examine which variables were associated with tau. A Pearson correlation coefficient was used to test correlations. A univariate linear regression was performed to examine which variables were associated with tau. A P value < 0.05 was considered as statistically significant.

Results

Anthropometric, clinical, and resting pulmonary function characteristics of both groups are listed in Table 1.

Spirometric evaluation in subjects with CF showed a mild obstructive pattern compared with healthy subjects with significantly reduced FEV₁ and FEV₁/FVC (Table 1), but without any statistically significant difference in IC between the 2 groups. Subjects with CF were significantly younger compared with healthy subjects (Table 1).

Table 2 summarizes the cardiorespiratory and metabolic exercise response of subjects with CF and healthy subjects during symptom-limited and constant work load submaximal CPET. There were no electrocardiogram findings indicating ischemia during symptom-limited CPET in either the CF or control group.

Exercise capacity was significantly reduced in subjects with CF compared with healthy subjects as assessed by VO₂peak and AT, whereas there was a significantly decreased O₂ pulse and an increased V̇E/V̇CO₂ slope. In the group of subjects with CF, there was significant prolonged oxygen recovery kinetics as expressed by VO₂/t slope compared with healthy subjects.

Subjects with CF had also a significantly increased duration of phase 2 VO₂ response during submaximal CPET as compared with healthy subjects (42.3 ± 21.5 vs 29.3 ± 6.4, s, P = .05) (Fig. 1). Phase 1 duration was prolonged in subjects with CF compared to healthy subjects; however, there was no statistically significant difference between the 2 groups.

A statistically significant negative correlation between FEV₁ (pred), FVC (pred), FEV₁/FVC, and tau was found (r = −0.77, P = .001, Fig. 2), (r = −0.79, P = .001), and (r = −0.62, P = .019), respectively in subjects with CF. Breathing reserve, VO₂peak (L/min), and AT (% VO₂peak) were also identified as significant correlates of tau (r = −0.74, P = .003, Fig. 3), (r = −0.53, P = .049, Fig. 4).
Fig. 1. Graphic representation of the time constant of the phase II response during constant workload exercise testing in adult subjects with cystic fibrosis and healthy controls. $P = .05$.

Fig. 2. Scatter graph of FEV1 (% predicted) vs time constant in adult subjects with cystic fibrosis.

Fig. 3. Scatter graph of breathing reserve vs time constant in adult subjects with cystic fibrosis.

Fig. 4. Scatter graph of $V_{O2peak}$ vs time constant in adult subjects with cystic fibrosis.

4), and [r = −0.65, $P = .013$]), respectively. $V_{O2}/t$ slope was significantly correlated with tau ($r = −0.58$, $P = .031$) in the same group. A statistically significant negative correlation between tau and Schwachman score was also found ($r = −0.80$, $P = .001$, Fig. 5).

A univariate linear regression was performed to examine which variables were significantly correlated to tau including FEV1(pred), FVC(pred), FEV1/FVC, $V_{O2peak}$ (L/min), breathing reserve, $V_E/V_{CO2}$ slope, and Schwachman score.

Multivariate linear regression analysis was conducted to determine the best linear combination of statistically significant univariate predictors for predicting tau. The results show that the Schwachman score ($β = −0.697$, $P = .002$) was the strongest independent predictor of tau ($R^2 = 0.719$, $F = 14.05$, $P = .001$).

Discussion

In the present study, we have shown that CF is associated with prolonged oxygen kinetics during submaximal constant work load exercise compared to healthy subjects. This oxygen kinetics prolongation was significantly correlated with disease severity as expressed by the common prognostic markers Schwachman score, FEV1, and $V_{O2peak}$ in CF.
SLOWED OXYGEN KINETICS IN CYSTIC FIBROSIS

$V_\text{O}_2$ kinetics of phase 2 of submaximal constant work load CPET have been used as an important index of central and peripheral limitation of exercise in various diseases such as COPD, heart failure, heart transplant recipients, cystic fibrosis, and even healthy subjects. The time constant of phase 2 oxygen kinetics (tau) is an index of aerobic metabolism of muscles and reflects aerobic capacity, but also is an index of oxygen delivery to peripheral tissues.

Until now, the data for oxygen kinetics of constant work load submaximal CPET in cystic fibrosis have been very limited. Hebestreit and co-workers have investigated the oxygen kinetics of constant work load submaximal CPET in subjects with CF and they have demonstrated that, although there was no difference in amplitude of phase 2 $V_\text{O}_2$ response between subjects with CF and healthy controls, the tau was prolonged. In the latter study, tau was adjusted only for oxygen desaturation and FEV1, leading to elimination of difference. Additionally, there was no documentation of the overall breathing pattern, and the CF population included children and adults (ages between 10 and 33 y old), facts that generate further queries.

Our data confirm prolongation of time constant tau in subjects with CF. Furthermore, our study has shown, for the first time, correlation of tau with disease severity as it is expressed by Schwachman score, FEV1, and peak exercise capacity. The inverse correlation between tau and indices of respiratory function at rest and during exercise (FEV1, FVC, FEV1/FVC, and breathing reserve) could indicate that phase 2 oxygen kinetics is significantly dependent on a central factor in CF and partly explains its prolongation by reduced oxygen delivery. This is in accordance with other studies in heart transplant recipients and COPD subjects. Most subjects with CF demonstrate an obstructive disease at spirometric evaluation. The prolonged exercise kinetics during maximal exercise has been partly explained by air trapping, ventilatory mismatch, muscle strength, nutritional status, and oxidative stress at the mitochondrial level. The prolongation of oxygen kinetics, which also emerged with submaximal constant work load exercise, indicates that the underlying mechanisms have early onset and probably affect overall clinical status and increase disease severity while decreasing exercise capacity. Moreover, indices of central limitation of exercise such as $O_2$ pulse and $V_\text{E}/V_\text{CO}_2$ slope were significantly different between the 2 groups. Subjects with CF exhibit a significantly lower $O_2$ pulse, an indirect index of stroke volume, and a significantly higher $V_\text{E}/V_\text{CO}_2$ slope, which is an index of ventilatory efficiency. These findings might indicate a reduced oxygen delivery, possibly due to some degree of pulmonary hypertension that is often present in CF, especially during exercise as has been reported elsewhere. However, we did not find any correlation between these indices of central limitation and phase 2 oxygen kinetics. Interestingly, cardiac function assessed by cardiac ultrasound at rest was within normal limits in our study, making this contribution to the prolonged oxygen kinetics in CF less possible. Moreover, although there was a slight increase in phase 1 duration in CF, this was not statistically significant. This phase reflects the cardiodynamic phase, which is rather normal in our adult subjects with CF.

In contrast to results in the study by Hebestreit et al, a statistically significant negative correlation was found between disease severity expressed by the Schwachman score and phase 2 oxygen kinetics. This correlation suggests that subjects with a high Schwachman score and consequently a good clinical condition demonstrate faster oxygen kinetics. To our knowledge, this is the first demonstration of a significant correlation between a clinical index of disease severity such as the Schwachman score and an index of aerobic capacity such as tau.

The inverse correlation of phase 2 oxygen kinetics with aerobic capacity ($V_\text{O}_2$AT) and early recovery oxygen kinetics after maximal exercise ($V_\text{O}_2$/t slope) in CF might also suggest that there is a peripheral factor that explains oxygen kinetics prolongation. Pouliou et al. have previously shown that there is prolonged early recovery oxygen kinetics after maximal exercise, which might indicate an impaired oxidative muscle metabolism of subjects with CF. Possible explanatory mechanisms for low oxidative capacity and prolonged oxygen kinetics in subjects with CF are deconditioning, abnormalities at the mitochondrial level such as increased calcium concentration, lower NADH dehydrogenase activity, and higher pH optimum of NADH dehydrogenase. Moreover, previous data suggest a reduced efficiency of oxidative ATP synthesis in subjects with CF. In the literature, an association between the main CF gene mutation and raised energy expenditure in subjects with CF is reported. These changes probably

Fig. 5. Scatter graph of Schwachman score vs time constant in adult subjects with cystic fibrosis.
affect muscle oxidative metabolism and may also explain phase 2 oxygen kinetics prolongation.

This study was cross-sectional, and not designed to prove causality. The study population consisted of a relative small sample size with healthy subjects not matched for age and BMI; however, the fact that, in the present study, subjects with CF were significantly younger than healthy subjects strengthens our findings. Younger subjects exhibit faster phase 2 oxygen kinetics.\(^3^6\) Subjects with CF even younger than healthy subjects exhibit slower phase 2 oxygen kinetics, outlining the role of central and peripheral factors to this phenomenon. The findings from our study are not valid for pediatric CF but only for adult CF. Moreover, the similar inspiratory capacities in both groups at baseline strengthens our data. Perpati et al\(^3^6\) have shown that lower baseline inspiratory capacity in subjects with CF correlates with poor exercise performance. The lack of echocardiographic data at the time of the study might limit our study conclusions; however, the negative medical history of cardiovascular risk factors for ischemic heart disease and a recent echocardiogram before referral at the CF outpatient clinic make cardiac involvement unlikely to explain the slow oxygen kinetics in these subjects. Definite explanatory mechanisms for prolonged oxygen kinetics during constant work load submaximal exercise cannot be deduced from the present findings; however, after thoroughly evaluating the symptom-limited CPET and its correlation with phase 2 oxygen kinetics, it seems that both central and peripheral factors influence prolonged oxygen kinetics. Further studies are needed to investigate the linking mechanisms.

**Clinical Implications**

These findings are clinically important, as the submaximal CPET is more tolerable and safer for debilitated subjects, who cannot perform maximal exercise. It could be used for risk stratification of subjects with CF. In the follow-up assessment, an accurate phase 2 oxygen kinetics evaluation instead of a symptom-limited CPET is suggested, avoiding the impact of low subject motivation and allowing physicians to terminate CPET considering both clinical and laboratory findings. Finally, submaximal exercise and tau estimation set the schedule of exercise training programs for subjects with CF and objectively assess the beneficial effects of rehabilitation.

**Conclusions**

Our study demonstrated that adult subjects with CF present a significant prolongation of oxygen kinetics during constant work load submaximal exercise testing in strong relation to disease severity. Both central and peripheral factors seem to be involved as explanatory mechanisms for this prolongation in subjects with CF.

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