Corticosteroids and skeletal muscle function in cystic fibrosis

Sinead C. Barry and Charles G. Gallagher

Department of Respiratory Medicine and the National Referral Centre for Adult Cystic Fibrosis, St. Vincent’s University Hospital, Dublin 4, Ireland

Submitted 11 June 2002; accepted in final form 30 May 2003

Barry, Sinead C., and Charles G. Gallagher. Corticosteroids and skeletal muscle function in cystic fibrosis. J Appl Physiol 95: 1379–1384, 2003. First published June 13, 2003; 10.1152/japplphysiol.00506.2002.—Patients with cystic fibrosis (CF) have reduced peripheral muscle strength. We tested the hypothesis that steroid treatment contributes to muscle weakness in adults with CF. Twenty-three stable CF patients were studied. Measurements included knee extensor (KE), knee flexor (KF), elbow flexor (EF), handgrip (HG), inspiratory (PImax), and expiratory (Pmax) muscle strengths. Spirometry, body mass index (BMI), and days spent in hospital over the preceding 12 mo (DH) were also measured. Average daily dose of prednisolone over the preceding 12 mo (ADD) was 5.1 mg/day. Pearson’s correlation analysis revealed that ADD correlated significantly with skeletal muscle strength (KE%, r = −0.63, P < 0.01) with the exception of HG%. These findings are independent of age, BMI, pulmonary function, and DH. Multiple-regression analysis revealed that ADD was the most significant predictor of all measures of skeletal muscle function except HG%. It was independently responsible for 54% of the variance in PImax%, for 46% of the variance in Pmax%, for 45% of the variance in KE%, for 39% of the variance in KE%, and for 41% of the variance in EF%. Concomitant medications (e.g., theophylline) were shown to have no causative effect. Corticosteroids contribute to the skeletal muscle weakness seen in CF patients. The correlation of proximal muscle strength, but not HG strength, with steroid dosage further supports a cause-effect relationship.

Many patients with cystic fibrosis (CF) have weak peripheral (9, 24) and, in some cases, respiratory muscles (23, 34). There is evidence that muscle weakness in CF may have major clinical implications (9, 16). Generalized muscle weakness may occur as a result of malnutrition (2, 14), disturbance in serum electrolytes (3), hypoxia (7, 8), and disuse (4, 15). Respiratory muscle weakness is often related to hyperinflation and poor nutrition (23, 34); however, preservation of respiratory strength has also been demonstrated in some CF patients despite the presence of low body weight and significant hyperinflation. Recent work by Decramer et al. (12) in chronic obstructive pulmonary disease has demonstrated a significant relationship between steroioid usage and both peripheral and respiratory muscle strength. The effect of steroid use on muscle strength in CF has not been examined to date. Despite the well-known negative side effects associated with chronic corticosteroid use, steroids still form an important part of the management of many patients with CF (5, 22, 33, 35).

We hypothesized that the muscle weakness seen in CF may be related in part to steroid use. We examined the relationship between skeletal muscle strength and steroid dosage (average daily dose in the previous year) in adults with CF who were clinically stable. We wanted to minimize the possibility that factors other than steroid usage might affect any relation between steroid dosage and muscle weakness. Therefore, we also examined the relation between skeletal muscle strength (dependent variable) and spirometry, nutritional status, and number of days spent in hospital over the previous year. Steroid myopathy preferentially affects proximal muscles and spares distal muscle groups. Therefore, we examined the relation between steroid dosage and strength of distal (e.g., handgrip) muscles as well as that between steroid dosage and strength of proximal muscles (e.g., knee flexor and extensor). We reasoned that if steroids contribute to muscle weakness in CF, proximal strength should correlate with steroid dosage but peripheral strength should not. Because acute infection or acute respiratory exacerbation may influence muscle function, patients were studied at least 1 mo after any acute exacerbation.

MATERIALS AND METHODS

Twenty-three patients (13 male, 10 female) were recruited from the outpatients department at the National Referral Centre for Adult Cystic Fibrosis at St. Vincent’s University Hospital, Dublin. Diagnosis of CF for each subject was based on clinical features, abnormal sweat test (sweat chloride > 60 mmol/l), and genotyping. All subjects were clinically stable for 1 mo before taking part in the study.

To assess the effects of corticosteroids on skeletal muscle function we measured inspiratory (PImax), expiratory (Pmax), knee extensor (KE), knee flexor (KF), elbow flexor (EF), and handgrip (HG) strengths. Pulmonary function tests were performed, and total steroid dose over the previous 12 mo was calculated in milligrams per day. All tests were performed during a single visit to the research laboratory. Be-
Table 1. Patient characteristics

<table>
<thead>
<tr>
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<th>Means ± SD</th>
<th>Range</th>
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</thead>
<tbody>
<tr>
<td>Age, yr</td>
<td>23.3 ± 5.1</td>
<td>18–39</td>
</tr>
<tr>
<td>BMI</td>
<td>20.6 ± 2.5</td>
<td>16.4–25.5</td>
</tr>
<tr>
<td>FEV1%</td>
<td>48.7 ± 24.0</td>
<td>25–108</td>
</tr>
<tr>
<td>FVC%</td>
<td>72.3 ± 20.6</td>
<td>33–112</td>
</tr>
<tr>
<td>DH</td>
<td>31.0 ± 30.9</td>
<td>0–100</td>
</tr>
</tbody>
</table>

Values are means ± SD for 23 patients (13 men, 10 women). FEV1%, forced expiratory volume in 1 s expressed as percentage predicted; FVC%, forced vital capacity expressed as percentage predicted; DH, days spent in hospital over the previous year; BMI, body mass index.

Table 2. Peripheral muscle strengths expressed as percentage predicted

<table>
<thead>
<tr>
<th></th>
<th>Means ± SD</th>
<th>Range</th>
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<tbody>
<tr>
<td>KE%</td>
<td>45.7 ± 17.6</td>
<td>15.8–91.3</td>
</tr>
<tr>
<td>KF%</td>
<td>62.0 ± 16.9</td>
<td>31.6–95.5</td>
</tr>
<tr>
<td>EF%</td>
<td>62.6 ± 18.1</td>
<td>24.3–105.0</td>
</tr>
<tr>
<td>HG%</td>
<td>67.9 ± 12.2</td>
<td>48.8–96.7</td>
</tr>
<tr>
<td>PEmax%</td>
<td>94.1 ± 34.2</td>
<td>39.5–175</td>
</tr>
<tr>
<td>Predmax%</td>
<td>102.4 ± 35.8</td>
<td>30.6–166.1</td>
</tr>
</tbody>
</table>

Values are means ± SD; n = 23. KE%, percentage predicted knee extensor strength; KF%, percentage predicted knee flexor strength; EF%, percentage predicted elbow flexor strength; HG%, percentage predicted handgrip strength; PEmax%, percentage predicted inspiratory muscle strength; Predmax%, percentage predicted expiratory muscle strength.
Prednisolone ADD was 5.1 mg/day. No subject was treated with corticosteroids for allergic bronchopulmonary aspergillosis.

Relationship between limb muscle and respiratory muscle strengths. Respiratory muscle strength and limb muscle strength correlated significantly with each other (P < 0.05); P\textsubscript{1max}% correlated with KE% (r = 0.76), KF% (r = 0.64), EK% (r = 0.87), HG% (r = 0.47), and P\textsubscript{max}% (r = 0.84). P\textsubscript{max}% was also significantly correlated with limb muscle strength, KE% (r = 0.73), KF% (r = 0.62), EK% (r = 0.83), and HG% (r = 0.51).

Relationship between muscle strength and steroid treatment. Using Pearson’s correlation analysis (Table 3; Fig. 1), we assessed the relationship between limb muscle and respiratory muscle strength and steroid dose and found a significant relationship between ADD and KE%, KF%, EF%, P\textsubscript{1max}%, and P\textsubscript{max}%. ADD and KE% and ADD and respiratory muscle strength (P\textsubscript{1max}%, P\textsubscript{max}%) were more significant than those seen between FEV\textsubscript{1}%, KE% and P\textsubscript{max}%. ADD explained 46% of the variance in P\textsubscript{max}%, with 22% being related to gender. For all muscle groups tested except HG%, ADD of corticosteroids explained the largest proportion of muscle weakness seen (Table 4). None of the concurrent medications contributed significantly to the regression model. Figure 3 illustrates the lack of correlation between theophylline dose expressed in milligrams per day and KE%.

DISCUSSION

The major new findings of this work are that 1) corticosteroid treatment independently correlates with limb and respiratory muscle strength in patients with CF; 2) the effects of corticosteroid treatment on skeletal muscle function are independent of airflow obstruc-
tion, nutrition, and DH; and 3) this correlation is seen for proximal muscles but not for the distal HG muscles. These results are in support of the hypothesis that treatments with corticosteroids have a negative effect on skeletal muscle strength in CF patients.

Possible limitations of this study are the following: 1) We used an indirect measurement of nutritional status, and no direct measure of lean body mass was performed. However, BMI has been shown to be a valuable measure of nutritional status in CF and to relate to exercise capacity (28), bone mineral density (17), and prognosis (20). 2) Our measures of limb muscle strength and respiratory muscle strength are effort dependent; however, each subject was tested repeatedly by the same investigator, making it difficult to perform repeated submaximal efforts. 3) We did not measure total body muscle mass, and as a result it is difficult to say conclusively whether we are seeing true myopathy, in which strength would be reduced at a disproportionate rate to muscle mass. 4) DH provides a useful marker of periods of extreme disuse; however, activity of daily living may contribute to disuse atrophy and has not been accounted for here.

A number of authors have looked at the presence of limb muscle weakness in CF patients. Lands et al. (24) demonstrated that leg strength was reduced in CF patients compared with controls and that strength was strongly related to body mass ($r = 0.816, P < 0.01$). Mier et al. (27) found that quadriceps strength was on average 68% predicted in their CF group. More recently, de Meer et al. (9) demonstrated that limb muscle force was reduced in CF children even in the absence of reduced pulmonary or nutritional status, whereas Elkin et al. (16) have demonstrated that both limb muscle strength and mass were reduced in their CF group compared with controls. The causes of muscle weakness in CF remain unclear; some authors suggest impairment in the quality of the muscle (9–11, 29), whereas others suggest that the differences in strength seen between CF patients and controls are related to body mass (16, 19, 24). No conclusive data are available at this time.

Studies of respiratory muscle function in CF are difficult to compare because of differences in the age of subjects studied, the methods used, and the indexes of muscle function. The tendency throughout the literature has been for inspiratory strength to be preserved in CF (20, 23, 24, 27, 30), and this is in agreement with our findings. Significant negative correlations were found between $P_{\text{Imax}}\%$, $P_{\text{Emax}}\%$, and ADD (Table 3). Our results demonstrate a wide range of values for $P_{\text{Imax}}\%$ (39.9–175%) with an equally wide range seen for $P_{\text{Emax}}\%$ (30.6–166.1%). We have shown that some patients have preserved respiratory muscle strength, whereas others have markedly reduced strength and still others have supernormal levels of respiratory muscle strength. The mechanisms responsible for these variations in strength are not clear. Factors such as reduced nutritional status, hypoxia, acidosis, electrolyte disturbances, and disuse may contribute to reduced strength, whereas chronic cough and increased work of breathing have been implicated in both preserved and increased respiratory muscle strength. It is likely that a number of factors simultaneously cause respiratory muscle variation. We have demonstrated a relationship between the widely varying respiratory strength seen in adults with CF and ADD, which have not been described previously.

There are many possible factors that can affect skeletal muscle function in CF. Weight loss, protein calorie insufficiency, abnormal ion transport, inflammation, sepsis, electrolyte disturbances, hypoxia, and corticosteroid usage all have myopathic potential. The purpose of this study was to assess the independent effects of corticosteroid therapy on limb muscle strength and respiratory muscle strength in a group of stable CF patients. It is known that myopathy may occur with virtually all steroids in common use (1, 13, 37). There are no previous data, to our knowledge, on the effects of treatment with corticosteroids on muscle strength in CF patients.

Like others, we have demonstrated the presence of limb muscle weakness in adults with CF (9, 16, 24, 27) and the preservation, on average, of respiratory muscle strength (21, 24, 30). We have shown that ADD of corticosteroids is significantly correlated with all measures of skeletal muscle function except HG%. We were unable to identify significant predictors of HG strength. HG was on average reduced at 67.9% predicted. Possible mechanisms of reduced HG strength are similar to those already described for generalized skeletal myopathy, with the exception of corticosteroid usage. These results are in agreement with our biological knowledge of steroid myopathy and its preference for proximal muscle atrophy, hence the lack of correlation with HG%.

Stepwise multiple-regression analysis revealed that ADD was responsible for the largest proportion of the muscle weakness seen in our CF patients, to a greater extent than nutritional status, pulmonary function, and DH. Single-regression analysis revealed a lack of correlation between peripheral muscle forces and $\text{FEV}_{1}\%$, BMI, age, and DH. The only significant corre-

![Fig. 3. Correlation of percentage predicted knee extensor strength with theophylline dose over the previous 12 mo.](image-url)
lation found was between KE% and FEV1%; however, this correlation was less than that seen for KE% and ADD. These findings suggest that treatment with corticosteroids contributes to muscle weakness in CF patients.

Nutritional status, disuse, and severity of disease all potentially affect muscle strengths in CF. To control for these potential cofactors, we assessed the effects of FEV1%, BMI, age, gender, and DH on strength of peripheral and respiratory muscles using both single- and multiple-regression analysis. Single-regression analysis revealed a significant relationship between KE% and FEV1%, which was less significant than between KE% and ADD. Stepwise multiple-regression analysis revealed that ADD was the most significant predictor of KE%, KE%, EF%, P1max%, and PEmax% with no significant effect on HG%. These results are in agreement with our hypothesis, i.e., that corticosteroids contribute to skeletal muscle weakness in CF.

In conclusion, this study demonstrates a significant association between corticosteroid dosage and skeletal muscle weakness in adults with CF. This correlation occurs at doses used in clinical practice. The correlation with steroid dosage is much greater than that attributable to age, sex, lung function, or overall nutritional status. These results have significant clinical implications.

DISCLOSURES

We thank the Cystic Fibrosis Research Trust, the Cystic Fibrosis Association of Ireland, the Health Research Board of Ireland, and the National Rehabilitation Board of Ireland for their support. We also thank the Cystic Fibrosis patients at St. Vincent’s University Hospital for their continued support.

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