Cystic fibrosis

Preferential reduction of quadriceps over respiratory muscle strength and bulk after lung transplantation for cystic fibrosis

C Pinet, P Scillia, M Cassart, M Lamotte, C Knoop, C Mélot, M Estenne

Background: In the absence of complications, recipients of lung transplants for cystic fibrosis have normal pulmonary function but the impact of the procedure on the strength and bulk of respiratory and limb muscles has not been studied.

Methods: Twelve stable patients who had undergone lung transplantation for cystic fibrosis 48 months earlier (range 8–95) and 12 normal subjects matched for age, height, and sex were studied. The following parameters were measured: standard lung function, peak oxygen uptake by cycle ergometry, diaphragm surface area by computed tomographic (CT) scanning, diaphragm and abdominal muscle thickness by ultrasonography, twitch transdiaphragmatic and gastric pressures, quadriceps isokinetic strength, and quadriceps cross section by CT scanning, and lean body mass. Diaphragm mass was computed from diaphragm surface area and thickness.

Results: Twitch transdiaphragmatic and gastric pressures, diaphragm mass, and abdominal muscle thickness were similar in the two groups but quadriceps strength and cross section were decreased by nearly 30% in the patients. Patients had preserved quadriceps strength per unit cross section but reduced quadriceps cross section per unit lean body mass. The cumulative dose of corticosteroids was an independent predictor of quadriceps atrophy. Peak oxygen uptake showed positive correlations with quadriceps strength and cross section in the two groups, but peak oxygen uptake per unit quadriceps strength or cross section was reduced in the patient group.

Conclusions: The diaphragm and abdominal muscles have preserved strength and bulk in patients transplanted for cystic fibrosis but the quadriceps is weak due to muscle atrophy. This atrophy is caused in part by corticosteroid therapy and correlates with the reduction in exercise capacity.
stable for the last 3 months before the study; (3) they were free of any decrease in forced expiratory volume in 1 second (FEV₁) suggesting the presence of chronic allograft dysfunction; and (4) they agreed to participate. At the start of the study in January 2002, 27 of the 52 patients with CF who had received a lung transplant at our institution since 1988 were alive. Of these 27 patients, 15 were excluded due to the presence of chronic rejection (n = 5), medical complications (n = 2), diaphragm paralysis (n = 1), and unwillingness to participate (n = 7), leaving 12 patients for study (five with HLT and seven with BLT). All patients but one were studied after the first postoperative year; the median time interval between transplantation and study was 47.5 months (range 8–95). At the time of the study the patients were taking a standard immunosuppressive regimen including cyclosporin (n = 5) or tacrolimus (n = 7), azathioprine (n = 8) or mycophenolate mofetyl (n = 1), and methylprednisolone (n = 12). Of the seven patients who were taking tacrolimus, five had been previously treated with cyclosporin for periods ranging from 22 days to 37 months.

The control group consisted of 12 normal subjects matched for age, height, and sex (table 1). These subjects, of whom 10 had participated in our previous study, were recruited from hospital personnel; they were all non-smokers and were free of any respiratory or neuromuscular disease.

All subjects were informed of the nature and extent of the study and gave written informed consent, as approved by the Human Studies Committee of the institution.

Pulmonary function
Measurements of standard pulmonary function were obtained by body plethysmography following the guidelines of the American Thoracic Society. Predicted values were computed using equations published by Quanjer et al. The patients performed an incrementally progressive, symptom limited, cardiopulmonary exercise test on an electrically braked cycle ergometer using standard techniques. After 5 minutes of adaptation to the mouthpiece the workload was increased by 10 W steps every minute to the point of exhaustion. The predicted value for peak oxygen uptake was derived from equations including sex, age, height, and weight.

Diaphragm mass
Diaphragm mass was computed from values of diaphragm surface area and thickness. Diaphragm surface area was measured using a previously described technique based on spiral computed tomographic (CT) scanning. The CT scanner (Somatom Volume Zoom, Siemens, AG, Erlangen, Germany) was also used to measure supine functional residual capacity (FRC). Diaphragm thickness was measured in the zone of apposition of the diaphragm to the lower rib cage by B-mode ultrasound using an 8–12 MHz linear probe (Toshiba Power Vision 8000, Japan), as previously described. In all subjects, measurements of diaphragm surface area and thickness were obtained during voluntary relaxation at FRC in the supine posture. The mean of three measurements of thickness was used for analysis.

Diaphragm strength
With the subject lying supine the two phrenic nerves were maximally stimulated transcutaneously in the neck at a frequency of 0.5 Hz with 0.2 ms square wave pulses delivered from two electrical stimulators (Medelec, Surrey, UK). The resulting twitch transdiaphragmatic pressure was measured using conventional balloon tipped catheters placed in the oesophagus and the stomach, as described elsewhere; the oesophageal and gastric balloons were filled with 0.5 and 1.0 ml of air, respectively. Subjects were instructed to breathe quietly for 20 minutes before stimulation to avoid twitch potentiation. Five runs of 5–6 twitches separated by 2–3 min of quiet breathing were obtained during voluntary relaxation against a closed airway at FRC; the last 2–3 runs were obtained during supramaximal stimulation. Twitches were considered valid and supramaximal based on standard criteria. The mean of the three twitches showing the largest transdiaphragmatic pressure was used for analysis.

Abdominal muscle thickness
The thickness of the rectus abdominis, external oblique, internal oblique, and transversus abdominis muscles was measured with a high resolution 8–12 Mhz ultrasound linear probe. Measurements were obtained on the right side with the subject relaxing at FRC in the supine posture, as previously described. Measurements were made in triplicate and the mean value was used for analysis. Abdominal muscle thickness was computed as the cumulative thickness of all four abdominal muscle layers.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Mean (SD) characteristics of study population</th>
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<tbody>
<tr>
<td></td>
<td>CF patients (n = 12)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>33.8 (6.6)</td>
</tr>
<tr>
<td>Sex</td>
<td>BM/4F</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>1.68 (0.07)</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>53.6 (8.5)</td>
</tr>
<tr>
<td>BMI (kg/㎡)</td>
<td>19.0 (1.8)</td>
</tr>
<tr>
<td>LBM (kg)</td>
<td>44.0 (8.5)</td>
</tr>
<tr>
<td>FBM (kg)</td>
<td>9.7 (3.6)</td>
</tr>
<tr>
<td>TLC (% pred)</td>
<td>100 (9)</td>
</tr>
<tr>
<td>FRC (% pred)</td>
<td>114 (20)</td>
</tr>
<tr>
<td>FEV₁ (% pred)</td>
<td>97 (14)</td>
</tr>
<tr>
<td>VC (% pred)</td>
<td>94 (12)</td>
</tr>
</tbody>
</table>

CF, cystic fibrosis; BMI, body mass index; LBM, lean body mass; FBM, fat body mass; TLC, total lung capacity; FRC, functional residual capacity; FEV₁, forced expiratory volume in 1 second; VC, vital capacity.

*Mean difference (with 95% confidence intervals) between CF and control groups.
measured and the mean of the three largest values was used for analysis.

**Quadriceps cross sectional area**
The cross sectional area of the quadriceps of the dominant leg was measured on a CT scan obtained midway between the femoral head and the medial femoral condyle. 11

**Quadriceps strength**
The maximum isokinetic (60°/s) strength of the quadriceps of the dominant leg was measured using a Cybex dynamometer.

**Nutritional status**
Nutritional status was assessed by computing body mass index, and by measuring lean and fat body mass using electrical bioimpedance (BIA, 101/S bioelectrical analyser, Akem, Florence, Italy). 21

**Data analysis**
The muscular surface area of the diaphragm (Amu) was calculated as Adi × 0.84 (where Adi is the total surface area of the muscle), based on the study by Arora and Rochester22 which showed that the surface area of the central tendon corresponds to 16% of total diaphragm surface area. Diaphragm mass was computed as Amu × Tdi × 1.04, where Tdi is diaphragm thickness and 1.04 is the density of the muscle. 23

Data are expressed as mean (SD) values or 95% confidence intervals (CI). Statistical analyses were made using paired and unpaired t tests, single linear regression analysis, and covariance analysis; the latter was used to test whether the vertical distance between the regression lines obtained in the two groups was significant. In addition, a multiple linear regression analysis was performed to investigate the inter-patient variability in the bulk and strength of the respiratory and quadriceps muscles. The following independent variables were included in the analysis: age, sex, body mass index, lean body mass, time between transplantation and study, type of transplant, presence of insulin dependent diabetes mellitus, mean daily dose of methylprednisolone received during the year preceding the study, and cumulative dose of methylprednisolone (calculated as the sum of all intravenous and oral doses received between surgery and study). The stepwise regression selection procedure was used to obtain the best model with the minimal number of predictors. Adjusted R² rather than R² values are reported to take into account the different numbers of predictors included in the models. A p value of <0.05 was considered statistically significant.

**RESULTS**
Mean anthropometric values and lung volumes in the patients and control subjects are shown in table 1. The two groups were matched for age, sex and height; weight, body mass index, fat body mass, and lean body mass tended to be smaller in the patients, but the differences did not reach statistical significance. Seven patients had insulin dependent diabetes mellitus. Mean values for pulmonary function tests in the patients were within normal limits; in particular, FRC was similar in the patients and the normal subjects, both in the seated and supine positions. The mean values for the bulk and strength of the diaphragm, abdominal, and quadriceps muscles are shown in table 2 and fig 1. Because of intolerance to oesophageal and gastric balloons, values of transdiaphragmatic and gastric pressure were not available in one patient; one control subject and one patient did not perform the exercise test. The mean twitch transdiaphragmatic pressure and diaphragm mass were similar in the patients and the normal subjects. In contrast, diaphragm mass normalised by lean body mass was 47% greater in patients than in controls (p = 0.002). Changes in gastric pressure elicited by abdominal muscle contraction were similar in the two groups. The cumulative thickness of the four abdominal muscle layers tended to be smaller in the patients than in the control subjects (p = 0.02), but this difference was exclusively caused by a reduction in the thickness of the external oblique and rectus abdominis muscles (p<0.02 and p = 0.005, respectively). The transversus abdominis and internal oblique muscles had a similar thickness in the two groups.

Quadriceps peak torque and cross sectional area were 33% (p = 0.006) and 31% (p = 0.001) lower in patients than in controls. Plots of peak torque versus cross sectional area showed positive correlations in both patients (r² = 0.77, p<0.001) and controls (r² = 0.75, p<0.001). Figure 2A shows that the relationships in the two groups were superimposed, indicating that quadriceps strength per unit cross sectional area was well preserved in the patients. On the other hand, mean values of cross sectional area per unit lean body mass were decreased by 20% in patients compared with controls (p = 0.001; fig 1, lower right panel). Plots of quadriceps cross sectional area versus lean body mass (fig 2B) showed significant correlations in the two groups (r² = 0.86, p<0.001 in controls and r² = 0.42, p<0.02 in patients), but the regression line computed in the patients was displaced towards lower values of muscle cross section (p = 0.002 by covariance analysis).

On average, peak oxygen uptake amounted to 56% of predicted (47–65%) and was not significantly different between patients receiving cyclosporin and those treated with tacrolimus. The reduced peak oxygen uptake was due to a reduced exercise time and power output, but the patients did not reach their maximal ventilation; on average, the ratio of minute ventilation at peak exercise over maximum voluntary ventilation amounted to 46% (37–55%). Nine patients stopped exercise

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Mean (SD) strength and bulk of respiratory and quadriceps muscles in the study population</th>
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<tbody>
<tr>
<td></td>
<td>CF patients</td>
</tr>
<tr>
<td>Pdi (cm H2O)</td>
<td>24.3 (4.5)</td>
</tr>
<tr>
<td>Mdi (g)</td>
<td>183 (53)</td>
</tr>
<tr>
<td>Mdi/LBM (g/kg)</td>
<td>4.22 (1.22)</td>
</tr>
<tr>
<td>Pga (cm H2O)</td>
<td>91.1 (28.8)</td>
</tr>
<tr>
<td>Tab (mm)</td>
<td>21.3 (4.7)</td>
</tr>
<tr>
<td>Tab/LBM (mm/kg)</td>
<td>0.49 (0.09)</td>
</tr>
<tr>
<td>PT (Nm)</td>
<td>103.6 (33.4)</td>
</tr>
<tr>
<td>quadCSA (cm²)</td>
<td>48.5 (13.8)</td>
</tr>
<tr>
<td>quadCSA/LBM (cm²/kg)</td>
<td>1.10 (0.26)</td>
</tr>
</tbody>
</table>

Pdi, transdiaphragmatic pressure elicited by twitch stimulation of the phrenic nerves; Mdi, diaphragm mass; LBM, lean body mass; Pga, change in gastric pressure elicited by abdominal stimulation; Tab, cumulative thickness of abdominal muscles; PT, peak torque of quadriceps muscle; quadCSA, cross sectional area of quadriceps muscle. *Mean difference between CF and control groups.
Figure 1  Upper panel: Mean values of transdiaphragmatic pressure (Pdi) elicited by twitch stimulation of the phrenic nerves in 11 CF patients and 12 controls, and of diaphragm mass (Mdi) in 12 CF patients and 12 control subjects. Middle panel: Mean changes in gastric pressure (Pga) elicited by stimulation of the abdominal muscles in 11 CF patients and 12 controls, and mean values of cumulated thickness of the abdominal muscles (Tab) in 12 CF patients and 12 control subjects. Lower panel: Mean values of quadriceps peak torque (PT) and cross sectional area (quad CSA) in 12 CF patients and 12 control subjects. LBM, lean body mass.

Figure 2  Relationships between (A) quadriceps peak torque (PT) and cross sectional area (quad CSA) and (B) quadriceps cross sectional area (quad CSA) and lean body mass (LBM) in 12 CF patients and 12 control subjects. The p value refers to the significance of the difference in vertical distance between the two regression lines as tested by analysis of covariance.
due to leg pain; two had no symptoms, but it was decided to stop the test because of a high respiratory quotient. For the 11 patients studied, the respiratory quotient at peak exercise averaged 1.31 (1.24–1.37).

Figure 3 shows that peak oxygen uptake was positively correlated with quadriceps strength ($r^2 = 0.78$ and 0.77, $p < 0.001$ in controls and patients, respectively) and cross sectional area ($r^2 = 0.73$, $p < 0.001$ and $r^2 = 0.68$, $p = 0.002$ in controls and patients, respectively). However, the regression lines computed in the patients were displaced towards lower values of peak oxygen uptake ($p = 0.007$ for quadriceps strength; $p = 0.07$ for cross sectional area by analysis of covariance). In accord with this, peak oxygen uptake per unit quadriceps strength was lower in patients (11.7 (1.9) ml/min/Nm) than in control (14.5 (2.2) ml/min/Nm; $p = 0.005$). The corresponding values for peak oxygen uptake per unit quadriceps cross sectional area were 24.7 (4.8) ml/min/cm$^2$ and 31.0 (4.4) ml/min/cm$^2$ ($p = 0.004$). Respiratory muscle strength and bulk did not correlate with peak oxygen uptake.

The mean daily dose of methylprednisolone taken by the patients during the year preceding the study amounted to 9.3 (4.3) mg and the total cumulative dose received between transplantation and study averaged 14.0 (5.6) g. The time elapsed since transplantation, the mean daily dose of methylprednisolone, and the cumulative dose of corticosteroids were not selected by multiple regression analysis as significant independent predictors of the bulk and strength of diaphragm and abdominal muscles, but they were predictive of quadriceps cross section according to the following equation: quadriceps cross section = $-5.64 + 1.26 \times \text{LBM} + 0.201 \times \text{time} - 0.0010 \times \text{Corticosteroids}$ (adjusted $R^2 = 0.83$, $p = 0.002$), where LBM is lean body mass, “time” is the time elapsed since surgery, and “corticosteroids” is the cumulative dose of methylprednisolone received since transplantation; replacing the latter by the mean daily dose of methylprednisolone did not improve the prediction. Quadriceps cross section thus tended to increase with postoperative time and to decrease with the total dose of corticosteroids. 43% of the variance in quadriceps cross section was due to LBM, 27% was related to the time elapsed since surgery, and the cumulative dose of corticosteroids was responsible for a further 13%. The strength and bulk of the respiratory and quadriceps muscles were similar in patients with HLT and BLT and in patients with and without diabetes mellitus.

**DISCUSSION**

This study has shown that, in patients transplanted for CF, the diaphragm and abdominal muscles have preserved strength and bulk but the quadriceps is weak due to muscle atrophy. This atrophy is caused partly by corticosteroid treatment and is closely correlated with the reduction in exercise capacity.

The patients and controls were not matched for weight because one aim of this study was to assess the impact of potential differences in nutritional status on the bulk of respiratory and quadriceps muscles. Body weight, BMI, lean body mass, and fat body mass were all non-significantly reduced in the patients compared with the controls, but analysis of the confidence intervals (table 1) indicates that the differences might have become significant if more patients had been included in the study (type II error). We did not quantify the level of daily physical activity in the two groups. The patients were included in a standard rehabilitation programme during the first 3–4 weeks after transplantation, but no formal revalidation was performed thereafter. At the time of the study two patients and three control subjects were engaged in a sports activity on a regular basis.

Previous studies of respiratory muscle strength in non-CF transplanted subjects have yielded conflicting results, possibly because of differences in patient characteristics in terms of time elapsed since surgery, underlying disease, and pulmonary function, and in the techniques used to assess muscle strength. In addition, interpretation of the data was made difficult by the small number of patients studied and the absence of a control group in most studies. This is the first study to use non-volitional techniques to show that the strength of the diaphragm and abdominal muscles is normal in patients with CF studied about 4 years after transplantation. Consistent with this finding, we found that the bulk of these muscles (normalised by lean body mass) was comparable to, or greater than, that measured in the control subjects. Furthermore, the mean daily dose of methylprednisolone and the cumulative dose of methylprednisolone taken by the patients were not selected by multiple regression analysis as independent predictors of respiratory muscle function and bulk. Together these observations indicate that corticosteroids given after transplantation do not produce significant respiratory muscle weakness or atrophy.

Several studies in non-transplanted CF patients have found limb muscle weakness which may be caused by several factors including inactivity (disuse atrophy), undernutrition, and chronic inflammation. Although these deleterious factors are all expected to improve after transplantation, the present study shows that quadriceps strength remained markedly decreased after the surgical procedure. This is consistent with a previous study which showed that...
leg strength was reduced to 76% of predicted in nine patients
with BLT (of whom seven had CF). Similarly, Ambrosino and
coworkers found a 35% decrease in quadriceps strength in
11 non-CF transplant recipients studied 1 year after
surgery. In the study by Pantoja and coworkers the strength
of the ankle dorsiflexor muscle measured in nine non-CF
patients 17 months after transplantation was decreased by
39%. Based on these reports, leg muscle weakness therefore
seems to be a consistent finding after lung transplantation
for both CF and other conditions.

Our study provides the novel and important finding that
this weakness is primarily related to muscle atrophy (rather
than myopathy). As shown in fig 2A, no patient had a
disproportionate decrease in strength compared with muscle
mass. Furthermore, the observation that quadriceps cross
section per unit lean body mass was significantly decreased
in patients compared with controls (fig 2B) indicates that
generalised muscle wasting caused by undernutrition could
not fully account for quadriceps atrophy—that is, there was
a disproportionate loss of quadriceps bulk. Although the
mechanisms underlying this alteration are unknown, the
toxicity of immunosuppressive agents is a likely contributor.
Based on previous reports, the mean daily dose of cortico-
steroids received by our patients seems sufficient to promote
muscle wasting, and the multiple regression analysis indicated
that the cumulative dose of methylprednisolone was an
independent predictor of quadriceps atrophy.

An intriguing finding of this study is that the long term use
of corticosteroids did not similarly result in atrophy of the
diaphragm and abdominal muscles. Acute myopathy with
weakness of respiratory and limb muscles induced by high
dose prednisolone has been reported in patients with a
variety of diseases including COPD or asthma and lung
transplantation, but the long term effects of administration
of a lower dose of corticosteroids is more controversial.
Some, but not all studies in chronic airflow
obstruction have shown that weakness of respiratory and/or
limb muscles is more likely to occur in patients receiving a
low dose of prednisolone. In patients with Cushing’s syn-
drome, Mills and coworkers found significant weakness of
the quadriceps but respiratory muscle strength was close to
normal. In fact, the available literature suggests that the
effects of corticosteroids on skeletal muscles may be
influenced by a variety of factors including the dose, duration
and type of exposure, type of steroids, type of muscle fiber,
and underlying disease. Corticosteroids may therefore be
more deleterious for muscles that are not continuously active,
are proximal, and contain a predominance of type IIb fibres,
features which all apply to the quadriceps of lung transplant
recipients. On this basis, it is possible to understand the
present finding that corticosteroids produced weakness and
wasting of the quadriceps but not of the respiratory muscles.

Seven of the 12 patients in our study had insulin
dependent diabetes mellitus which may cause polynu-
ereathy and muscular atrophy. Based on the present findings,
we cannot precisely determine the extent to which diabetes
contributed to quadriceps wasting, but the following obser-
vations argue against an important role. Firstly, polynu-
ereathy is more likely to occur in patients with longstanding
and/or ill controlled diabetes. In our patients the median
duration of diabetes was 24 months and only one patient had
had diabetes for more than 5 years; in addition, the more
recent value for glycated haemoglobin at the time of study
averaged 7.5%, indicating fairly good control of the disease.
Secondly, quadriceps strength and cross section did not differ
in patients with and without diabetes. Finally, the presence
of diabetes was not selected by the multiple regression
analysis as a significant independent predictor of quadriceps
atrophy.

Some of the patients were treated with cyclosporin which
may also have adverse effects on skeletal muscles. A number
of case studies have reported muscular disorders
associated with cyclosporin, including myopathy, but it was
generally difficult to exclude a causative or potentiating role
for other medications such as corticosteroids or statins. In a
study in rats, Biring and coworkers showed that cyclosporin
alone had no effect on the proportion and cross sectional area
of diaphragm and limb muscle fibres. A direct effect of
cyclosporin on the bulk of the quadriceps therefore seems
unlikely. To the best of our knowledge, the potential toxicity
of tacrolimus on skeletal muscle has not been studied to date.

Our analysis of the relationships between peak oxygen
uptake and quadriceps strength or cross sectional area pro-
vides some insight into the factors limiting exercise capacity
in transplanted CF patients. We observed that the relation-
ships obtained in the patients were displaced towards lower
values of peak oxygen uptake (fig 3)—that is, mean values of
peak oxygen uptake per unit quadriceps strength or cross
sectional area were lower in patients than in controls. These
observations indicate that quadriceps atrophy alone could
not account for all the impairment in peak oxygen uptake.
Several studies in lung transplant recipients have indicated a
persistent impairment in exercise capacity, which is present
independent of the type of transplant procedure and involves
recipients with and without CF. This impairment has been
attributed to a defect originating in the peripheral muscles
and limiting oxygen utilisation; this defect may be caused by
a generalised deconditioning and by the toxicity of cyclo-
sporin which decreases the oxidative capacity of peripheral
muscles by inhibiting mitochondrial respiration and decreas-
ing capillary density. Other possible contributors to the
disproportionate decrease in peak oxygen uptake in compari-
son with quadriceps strength and bulk include a CF related
abnormality in muscle oxygen metabolism, increased
sensation of fatigue in the peripheral muscles, or impaired
oxygen delivery to these muscles, particularly in recipients of
heart-lung transplants.

Although the present data do not allow us to assess the
relative role played by quadriceps wasting and by derange-
ments in muscle oxygen metabolism, it seems very likely that
restoring muscle mass would improve exercise tolerance in
transplanted CF patients. Previous studies using aerobic
endurance training alone in lung transplant recipients have
shown only modest improvements in maximal oxygen
uptake after 11 weeks of training (for example, +11.5% com-
pared with baseline). It is likely that adding strategies
aimed at increasing muscle bulk such as strength training or
anabolic drugs to endurance training alone will produce
greater gains in exercise tolerance.

In summary, this cross sectional study has shown that the
strength and bulk of the diaphragm and abdominal muscles
is preserved in patients who have undergone lung transplan-
tation for CF while the strength of the quadriceps muscle
is markedly decreased. Our results also indicate that this
reduction is entirely accounted for by muscle atrophy. This
atrophy is caused in part by corticosteroid therapy and is
considerably correlated with the reduction in exercise capacity.
Longitudinal studies of limb muscle function, bulk, and
histology before and after surgery are now required to acquire
a better understanding of the effects of lung transplantation
on skeletal muscles and of the mechanisms responsible for
quadriceps atrophy.

ACKNOWLEDGEMENTS

The authors thank B Morlion, R C Sà, and M Paiva (Biomedical
Physic Laboratory, ULB, Brussels, Belgium) for their support in the
data analysis and G Gayan-Ramirez (Respiratory Muscles Laboratory,
KUL, Leuven, Belgium) for helpful discussions on the data.
The study was supported by the Association Belge de Lutte contre la Mucoviscidose (ABLM).

REFERENCES


