Original articles

Wasting as an independent predictor of mortality in patients with cystic fibrosis

R Sharma, V G Florea, A P Bolger, W Doehner, N D Florea, A J S Coats, M E Hodson, S D Anker, M Y Henein

Abstract

Background—Cystic fibrosis (CF) is the most common life threatening autosomal recessive disorder in the white population. Wasting has long been recognised as a poor prognostic marker in CF. Whether it predicts survival independently of lung function and arterial blood gas tensions has not previously been reported.

Methods—584 patients with CF (261 women) of mean (SD) age 21 (7) years were studied between 1985 and 1996, all of whom were being followed up in a tertiary referral centre. Lung function tests, body weight, arterial blood oxygen (PaO₂) and carbon dioxide (PaCO₂) tensions were measured. The weight was calculated as a percentage of the ideal body weight for age, height, and sex.

Results—Forced expiratory volume in one second (FEV₁) recorded at the start of the study was 1.8 (1.0) l (52 (26)% predicted FEV₁), PaO₂ 9.8 (1.9) kPa, PaCO₂ 5.0 (0.9) kPa, and % ideal weight 92 (18)%.

During the follow up period (45 (27) months) 137 patients died (5 year survival 72%, 95% CI 67 to 73). FEV₁, % predicted FEV₁, PaO₂, % ideal weight (all p<0.0001), and PaCO₂ (p=0.04) predicted survival. In multivariate analysis, % predicted FEV₁ (p<0.0001), % ideal weight (p=0.004), and PaCO₂ (p=0.02) were independent predictors of outcome. Patients with >85% ideal body weight had a better prognosis at 5 years (cumulative survival 84%, 95% CI 79 to 89) than those with ≤85% ideal weight (survival 53%, 95% CI 45 to 62), p<0.0001.

Percentage predicted FEV₁ (area under curve 0.83; 95% CI 0.78 to 0.87) and % ideal weight (area under curve 0.74; 95% CI 0.68 to 0.79) were accurate predictors of survival at 5 years follow up (receiver-operating characteristic analysis).

Conclusions—Body wasting is a significant predictor of survival in patients with CF independent of lung function, arterial blood oxygen and carbon dioxide tensions.

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Keywords: cystic fibrosis; wasting; survival

Cystic fibrosis (CF) is the most common life threatening autosomal recessive disorder in the white population, occurring at a frequency of approximately 1:2000 to 1:2500 live births. Advances in medical treatment have resulted in an improvement in prognosis and symptoms with increasing numbers of patients surviving into adult life. However, as the disease progresses, patients develop disabling lung disease with the main cause of death in adults being respiratory failure. For patients with advanced CF, lung or heart-lung transplantation represents an important treatment for improving survival and quality of life. In view of the shortage of donor organs and to assess the suitability of patients for transplantation, reliable prognostic indicators that can accurately define groups of patients at high risk are needed.

Wasting has long been recognised as a marker of impaired survival in CF, but it has not been previously shown to be independent of established prognostic indicators. Kerem et al. showed that body wasting was predictive of impaired survival but this was not independent of lung function or arterial blood gas tensions. It has been demonstrated that cachexia is a strong and independent risk factor for mortality in patients with chronic heart failure, as well as other chronic diseases. We have examined whether body wasting is an independent prognostic marker in CF.

Methods

STUDY POPULATION AND FOLLOW UP

Between 1985 and 1996, 584 patients (261 female) with CF, of mean age 21 (7) years (range 7–45), who attended the Royal Brompton Hospital were studied. The diagnosis of CF

Table 1 Baseline clinical characteristics of 584 consecutive patients with cystic fibrosis (CF)

<table>
<thead>
<tr>
<th>All CF patients (n=584)</th>
<th>Died (n=137)</th>
<th>Alive (n=447)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>21 (7)</td>
<td>22 (7)</td>
</tr>
<tr>
<td>Sex (MF)</td>
<td>323/261</td>
<td>73/64</td>
</tr>
<tr>
<td>FEV₁ (l)</td>
<td>1.8 (1.0)</td>
<td>1.1 (0.5)</td>
</tr>
<tr>
<td>% predicted FEV₁ (%)</td>
<td>52 (26)</td>
<td>32 (15)</td>
</tr>
<tr>
<td>PaO₂ (kPa)</td>
<td>9.8 (1.9)</td>
<td>8.7 (1.5)</td>
</tr>
<tr>
<td>PaCO₂ (kPa)</td>
<td>5.0 (0.9)</td>
<td>5.3 (1.1)</td>
</tr>
</tbody>
</table>

Data are presented as mean (SD) or numbers of patients.

FEV₁ = forced expiratory volume in one second; % ideal weight = weight as percentage of the predicted ideal weight; PaO₂ = arterial oxygen tension; PaCO₂ = arterial carbon dioxide tension.

*p<0.01, †p<0.0001 for unpaired t test of mean of corresponding groups.

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FEV1 = forced expiratory volume in one second; PaO2 = arterial oxygen tension; PaCO2 = arterial carbon dioxide tension; % ideal weight = weight as percentage of the predicted ideal weight.

**Table 2** Cox proportional hazards analysis of survival in 584 patients with cystic fibrosis: univariate analysis

<table>
<thead>
<tr>
<th>Parameter</th>
<th>$\chi^2$ value</th>
<th>p value</th>
<th>RR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>FEV1</td>
<td>137.0</td>
<td>&lt;0.0001</td>
<td>0.999 (0.998 to 0.999)</td>
</tr>
<tr>
<td>% predicted FEV1</td>
<td>143.3</td>
<td>&lt;0.0001</td>
<td>0.945 (0.934 to 0.956)</td>
</tr>
<tr>
<td>% ideal weight</td>
<td>62.0</td>
<td>&lt;0.0001</td>
<td>0.995 (0.942 to 0.957)</td>
</tr>
<tr>
<td>PaO2</td>
<td>21.7</td>
<td>&lt;0.0001</td>
<td>0.730 (0.637 to 0.838)</td>
</tr>
<tr>
<td>PaCO2</td>
<td>4.0</td>
<td>0.04</td>
<td>1.280 (1.009 to 1.622)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>0.7</td>
<td>0.406</td>
<td>1.010 (0.986 to 1.034)</td>
</tr>
<tr>
<td>Female sex</td>
<td>1.2</td>
<td>0.280</td>
<td>1.203 (0.860 to 1.684)</td>
</tr>
</tbody>
</table>

FEV1 = forced expiratory volume in one second; PaO2 = arterial oxygen tension; PaCO2 = arterial carbon dioxide tension; % ideal weight = weight as percentage of the predicted ideal weight.

**Table 3** Cox proportional hazards analysis of survival in 584 patients with cystic fibrosis: multivariate analysis

<table>
<thead>
<tr>
<th>Parameter</th>
<th>$\chi^2$ value</th>
<th>p value</th>
<th>RR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>% ideal weight</td>
<td>5.9</td>
<td>0.015</td>
<td>0.984 (0.971 to 0.997)</td>
</tr>
<tr>
<td>% predicted FEV1</td>
<td>63.4</td>
<td>&lt;0.0001</td>
<td>0.951 (0.940 to 0.963)</td>
</tr>
<tr>
<td>% ideal weight</td>
<td>56.5</td>
<td>&lt;0.0001</td>
<td>0.955 (0.944 to 0.967)</td>
</tr>
<tr>
<td>PaCO2</td>
<td>4.0</td>
<td>0.04</td>
<td>1.280 (1.009 to 1.622)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>0.7</td>
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<tr>
<td>Female sex</td>
<td>1.2</td>
<td>0.280</td>
<td>1.203 (0.860 to 1.684)</td>
</tr>
</tbody>
</table>

FEV1 = forced expiratory volume in one second; PaO2 = arterial oxygen tension; PaCO2 = arterial carbon dioxide tension; % ideal weight = weight as percentage of the predicted ideal weight.

**Figure 1** Kaplan-Meier survival plot for 584 patients with cystic fibrosis: patients subgrouped according to (A) percentage ideal weight and (B) percentage predicted forced expiratory volume in one second (FEV1).

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was based on positive sweat tests with typical clinical findings, with or without genotype confirmation. The patients underwent routine testing of pulmonary function during their follow up, with forced expiratory volume in one second (FEV1) expressed as a percentage of the normal predicted values for height and sex (% predicted FEV1), as previously described. 9 10 Height, weight, arterial blood oxygen and carbon dioxide tensions (PaO2, PaCO2) were also measured. Weight was calculated as a percentage of the ideal body weight for height, age, and sex using the standards of Tanner 11 for patients aged ≤19 years and the Metropolitan height and weight tables12 for patients >19 years of age.

As all the above investigations were part of the routine assessment of patients with CF attending our hospital, informed consent from the patients studied or ethics committee approval was deemed unnecessary.

**STATISTICAL ANALYSIS**

All results are presented as mean (SD) values. Cox proportional hazard analysis was performed using baseline values to assess the association between variables and all-cause mortality. Previous studies have shown that % predicted FEV1, age, and sex all relate to survival. The interaction between the effects of each of these variables and % ideal weight in determining survival was therefore evaluated initially in two variable models. Subsequently, these parameters (together with arterial blood gas levels) were included in a six variable model (table 3). Hazard ratio (RR) and 95% confidence interval (CI) for risk factors as well as significance levels for $\chi^2$ (likelihood ratio test) are given, and Kaplan-Meier cumulative survival plots were constructed (StatView 5, Abacus Concepts, Berkeley, USA). The receiver-operating characteristic curves for % predicted FEV1, % ideal weight, and PaCO2 were drawn and the areas under the curves calculated (MedCalc 5.0, MedCalc Inc, Belgium). The cut off level that resulted in the highest product of sensitivity and specificity was considered the optimal level for predicting outcome.

The probability of death within 5 years of patients with CF (according to % predicted FEV1 and % ideal weight) was calculated using a logistic regression analysis technique. 13

**Results**

Of the 584 patients who were followed up for a mean of 45 (27) months, 137 (23.5%) died after 2.4–89.9 months (mean 31.0 (21.0); median 25.0). The mean (SD) follow up period of the 447 survivors was 48.0 (27.1) months (range 1.6–94.7). The cumulative survival of all patients was 96.3% at 1 year (95% CI 94.7 to 97.9), 82.8% at 3 years (95% CI 79.4 to 86.2), 71.9% at 5 years (95% CI 67.3 to 76.5), and 62.7% at 7 years (95% CI 56.9 to 68.5). For all the 584 patients the mean FEV1 recorded at the start of study was 1.8 (1.0) l (52 (26%) predicted FEV1), PaO2 9.8 (1.9) kPa, PaCO2 5.0 (0.9) kPa, and % ideal weight 92 (18)% (table 1).
Receiver-operating characteristic (ROC) curves were plotted for % predicted FEV₁, % ideal weight, and PaCO₂ (fig 2). Both % predicted FEV₁ (area under the curve 0.827; 95% CI 0.780 to 0.867) and % ideal weight (area under the curve 0.736; 95% CI 0.684 to 0.785) were found to be highly accurate predictors of 5 year survival, with the former being significantly better than the latter (p=0.002). In contrast, PaCO₂ was not shown to be an accurate prognostic indicator (area under the curve 0.543; 95% CI 0.445 to 0.639). ROC curves were also drawn for 1, 2, 3, and 7 years follow up which resulted in similar results to those above (data not shown). The optimal cut off value for predicting death at 5 years was 38% for predicted FEV₁, and 83% for ideal weight.

The probability of death within 5 years was estimated according to % predicted FEV₁ and % ideal weight (fig 3). For any given % predicted FEV₁, the probability of death was found to increase markedly as the % ideal weight decreased. The probability of death within 2 and 7 years was also calculated, resulting in similar findings (data not shown).

**Discussion**

This study has shown that body wasting is an important predictor of impaired survival in CF. It provides prognostic information independently of established risk factors such as lung function (% predicted FEV₁) and arterial blood gas tensions (PaO₂ and PaCO₂). Although it has long been recognised that wasting is an independent predictor of survival in CF, it has not been previously shown to be an independent prognostic indicator.

In a study by Kerem et al, % ideal weight was not found to be an independent predictor of survival, in contrast to the present study. One reason for this may be that our data are more recent (patients followed up between 1985 and 1996) than those of Kerem et al (patients followed up between 1977 and 1989). During the last two decades there have been major improvements in the nutrition of patients with CF, with the introduction of high calorie diets and the development of better pancreatic enzymes in the mid 1980s. It may be hypothesised that, in the past, if a patient with CF was cachectic this was simply due to poor nutrition, which is why body weight was a relatively weak marker of disease severity in the study by Kerem et al. However, since the advances in nutritional support, the development of weight loss (or failure to thrive in children) is likely to be due to a serious deterioration in the metabolic status of the patient as a result of the disease process itself. As a result, a low % ideal body weight has become a more powerful and independent predictor of survival in patients with CF who are managed using present day treatment strategies. It must be borne in mind that, as new therapies for chronic diseases emerge, the prognostic value of certain markers may change, as has been observed in other disease states.

In the present study, age and sex were not significant predictors of mortality in CF.
There are several reasons for weight loss in patients with CF, including uncontrolled malabsorption, the development of diabetes mellitus, or the progression of respiratory disease. Recurrent episodes of pulmonary infection can result in wasting due to an increase in metabolic rate and because the patient may become too breathless to eat properly. Cachexia is also known to be a poor prognostic sign in patients with other medical conditions such as cancer, AIDS, and chronic heart failure.4 5 Although there are different mechanisms responsible for the wasting process in these conditions, the common finding appears to be that cachexia is a strong predictor of impaired survival. Furthermore, it is known that improvements in nutrition can lead to increased survival of patients with CF. In a study comparing survival rates of two large CF centres in North America, there was a higher median survival rate among patients in Toronto than in those in Boston because of better nutrition in the former centre.22

The present study shows that CF patients with wasting represent a high risk group which should be considered for transplantation at an earlier stage than patients who are not cachectic. Previous work has shown that preoperative body mass index does not influence survival following transplantation, suggesting that cachectic patients should not be denied transplantation unduly while attempts are being made to increase weight.17 Percentage ideal weight represents a simple and cost effective measurement which, in addition to % predicted FEV1, provides an accurate and independent prediction of prognosis in CF.

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1 Hodson ME. Cystic fibrosis. Postgrad Med J 1984;60:225–33.