Lung clearance index predicts pulmonary exacerbations in young patients with cystic fibrosis

François Vermeulen, Marijke Proesmans, Mieke Boon, Trudy Havermans, Kris De Boeck

ABSTRACT

Rationale The lung clearance index (LCI) is a promising endpoint for use in cystic fibrosis (CF) clinical trials, but correlations with validated clinical endpoints have not yet been established.

Objective This study aimed to demonstrate that, in young patients with CF, baseline LCI predicts subsequent pulmonary exacerbation (PE) and correlates with the respiratory domain of the CF Questionnaire-Revised (CFQ-Rresp).

Methods Baseline LCI, forced expiratory volume in 1 s (FEV1), CFQ-Rresp and PEs over the subsequent year were prospectively recorded in 63 patients aged 5–19 years. The ability of baseline LCI to predict PE was assessed using negative binomial regression models and Kaplan–Meier plots.

Results Twenty-six patients (41%) experienced 48 PEs. Baseline LCI and FEV1 were predictors of PE. Compared with the quartile with the lowest LCI, the annual PE rate in increasing LCI quartiles was 2.9 (95% CI 0.5 to 16.5, p=0.238), 5.4 (95% CI 1.0 to 29.0, p=0.045) and 13.6 (95% CI 2.8 to 67.1, p=0.001). Similarly, time to first PE decreased with worsening LCI quartiles (log-rank test for trend, p<0.001). Furthermore, LCI correlated with CFQ-Rresp (r=−0.43, p<0.001). In the subgroup of 53 patients with normal FEV1, LCI was a predictor of PE. In this subgroup, LCI also correlated with CFQ-Rresp (r=−0.282, p=0.043).

Conclusions Baseline LCI predicts PE in young patients with CF and correlates with CFQ-Rresp, a validated patient-reported outcome, even in the subgroup with normal FEV1. These data further support the use of LCI as a surrogate outcome measure in CF clinical trials.

INTRODUCTION

Forced expiratory volume in 1 s (FEV1) is currently the only accepted surrogate endpoint for use in cystic fibrosis (CF) clinical trials. As most patients with CF have a normal FEV1 until adolescence, FEV1 is an insensitive endpoint for this age group. New surrogate endpoints are therefore needed. The lung clearance index (LCI), an index of ventilation homogeneity derived from the multiple breath washout (MBW) of an inert gas, is one candidate surrogate endpoint, as is chest CT.

The LCI satisfies many of the conditions needed to qualify as a surrogate endpoint. It is feasible in a ‘standard’ clinical setting and can be measured repeatedly without harm. The measurement technique was recently standardised and commercially available devices have been validated. Furthermore, the short- and medium-term repeatability are good in both healthy subjects and patients, and the LCI is more sensitive than FEV1 for detecting early CF lung disease in cross-sectional analyses. The LCI is also responsive to interventions such as treatments with intravenous antibiotics, hypertonic saline or rh-DNase, and it was used successfully as an endpoint in a trial evaluating the effect of hypertonic saline in infants. In addition, the LCI was shown to be improved in young children with mild CF lung disease carrying at least one G551D mutation who were treated with ivacaftor. Initial data concerning the correlation between the LCI and long-term disease course have emerged, and abnormal LCI values in preschool children with CF were shown to predict spirometric abnormalities at school age.

To validate a physiological measurement as a surrogate endpoint it should be demonstrated that it is correlated with validated clinical outcomes such as survival or, in the case of CF, pulmonary exacerbations (PEs) or patient-reported outcomes has to be established.

Key messages

What is the key question?

The lung clearance index (LCI) is a marker of early cystic fibrosis (CF) lung disease proven to be sensitive, repeatable and responsive to interventions. However, before the LCI can be used as a surrogate endpoint in trials involving patients with CF, a correlation with clinically meaningful endpoints such as pulmonary exacerbation or patient-reported outcomes has to be established.

What is the bottom line?

In young patients with CF, baseline LCI predicts the time to the first pulmonary exacerbation in the 12 months after baseline assessment. The LCI also correlates with CFQ-Rresp, a validated patient-reported outcome.

Why read on?

This prospective work demonstrates the link between the LCI and two clinically meaningful endpoints in young patients with CF: pulmonary exacerbations and CFQ-Rresp. This contributes to the validation of the LCI as a surrogate endpoint for use in clinical trials.
subsequent PE, patient-reported outcome measures and symptoms as a further step in the validation of the LCI as a surrogate endpoint for clinical trials in patients with CF.

Our first hypothesis was that the baseline LCI would predict subsequent PEs during the year after the baseline measurements. Our second hypothesis was that the LCI would correlate with patient-reported outcome measures and symptom scores. Because the LCI is more sensitive than FEV₁ for detecting early lung disease in CF, the LCI was expected to be a predictor of PE and to correlate with patient-reported outcome measures even in the subgroup of patients with normal FEV₁.

Some of the results of this study have been previously reported in the form of an abstract.¹⁶

METHODS

Recruitment
In this prospective single-centre study, patients with CF aged between 5 and 20 years were recruited during routine outpatient visits between January 2011 and October 2012. Patients infected with methicillin-resistant *Staphylococcus aureus* (MRSA) or *Burkholderia cepacia* are not allowed in the standard lung function laboratory and were excluded.

Control subjects, recruited by advertisement, had no history of chronic respiratory symptoms and had not suffered respiratory infections in the previous 4 weeks.

The study was approved by the ethics committee of the University of Leuven. Caregivers and/or patients gave written informed consent before inclusion.

Multiple breath washout (MBW) measurement

A detailed description of MBW measurements and quality control is shown in the online supplement.

Nitrogen MBW was performed using the Exhalyzer D (EcoMedics AG, Duernten, Switzerland), as described elsewhere.³ Before spirometry and after inhalation of 400 µg salbutamol, MBW measurements were performed with a between-test delay of more than twice the washout time of the previous measurement.

LCI calculations and repeatability criteria

An MBW measurement was excluded if significant leaks, sighs or irregular breathing occurred or if the functional residual capacity (FRC) differed by more than 20% compared with the largest FRC of the valid measurements within the same session. The mean LCI of all valid MBW measurements was used.

Z-scores were calculated from the values in the control group. An LCI z-score of >2 was considered abnormal.

Spirometry and biometry

FEV₁, forced vital capacity (FVC) and forced expiratory flow between 25% and 75% of FVC (FEF₂₅–₇₅) were expressed as the percent predicted or as z-scores using reference equations from the Global Lung Function Initiative.¹⁷ A z-score lower than –2 was considered abnormal.

Height, weight and body mass index were expressed as z-scores according to local reference equations.¹⁸

Pulmonary exacerbations

The time to the first PE and the number of PEs during the 12 months following the baseline LCI measurement were recorded. PEs were defined as changes in respiratory status for which intravenous antibiotics were administered. Intravenous antibiotic therapy was prescribed by the CF consultant in accordance with current practice guidelines.¹⁷ The LCI results were not disclosed to the clinicians.

Patient-reported outcome and symptom score

Before any other procedure, each child (≥12 years) or parent (<12 years) completed the respiratory domain of the CF Questionnaire-Revised (CFQ-R resp).²⁰ The CFQ-R resp was standardised to range from 0 to 100, with high scores indicating fewer respiratory symptoms.

The CF Clinical Score (CFCS)¹⁰ is a composite score including symptoms and clinical findings assessed by the CF consultant and ranges from 10 (mildest symptoms and signs) to 50 (most severe symptoms and signs).

Both assessments were taken on the same day as the MBW measurement and the spirometry.

Statistical analysis

The LCI, spirometric parameters, CFQ-R resp and CFCS were divided into quartiles. The time to subsequent PE was compared between quartiles using Kaplan–Meier plots and the log-rank test for trend.

The predictive value of age, gender, infection with *Pseudomonas aeruginosa*, LCI z-score, FEV₁ z-score, FEF₂₅–₇₅ z-score, CFQ-R resp and CFCS on annualised PE rate was evaluated using univariate and multivariate negative binomial regression models.

Correlations between parameters were assessed by Spearman correlation coefficient (r). SPSS V21.0 (IBM, Armonk, New York, USA) was used for the statistical analyses.

RESULTS

Study population

Of the 86 eligible 5–19-year-old patients with CF in follow-up at the Leuven CF Centre, 63 were enrolled. The reasons for not participating included patient not approached by the staff (n=13), infection with *B cepacia* and/or MRSA (n=5), inability to obtain a valid MBW measurement (n=3) or refusal by the patient (n=2). There were no differences in age, weight, height, body mass index, spirometry, pancreatic status, proportion of F508del homozygotes or colonisation status for *P aeruginosa* between patients who were or were not enrolled (table 1).

Of the 61 enrolled control subjects, three were excluded due to inability to perform the MBW measurement and one because of suspicion of subclinical lung disease (Ehler–Danlos syndrome). Compared with the patients with CF, the control subjects had a similar age distribution but a greater z-score for weight, height, FEV₁ and FEF₂₅–₇₅ and a lower LCI z-score (table 1).

Correlation between LCI, FEV₁ and FEF₂₅–₇₅

In the patients with CF there was a negative correlation between the LCI z-score and FEV₁ z-score (r=–0.642, p<0.001). In 42 of the 63 patients (67%) the LCI was abnormal but the FEV₁ was normal. Eleven patients (17%) had a normal LCI and a normal FEV₁ and 10 (16%) had an abnormal LCI and FEV₁. None of the patients with an abnormal FEV₁ had a normal LCI (figure 1 and online supplementary figure E1). There was a moderate correlation between the LCI and FEF₂₅–₇₅ z-scores (r=–0.673, p<0.001, see online supplementary figure E2) and a strong correlation between FEV₁ and FEF₂₅–₇₅ z-scores (r=0.731, p<0.001). Of 53 patients with a normal FEV₁, 47 (89%) also had a normal FEF₂₅–₇₅ and 6 (11%) had an abnormal FEF₂₅–₇₅. These six patients also had an abnormal LCI (see online supplementary figure E2).
All control subjects had a normal LCI and a normal FEV₁. A negative correlation between the LCI z-score and FEV₁ z-score was also observed in the controls (r = −0.384, p = 0.004).

Correlation of lung function with CFQ-Rresp and CFCS

The LCI z-score (r = −0.431, p < 0.001, figure 2A) and the FEV₁ z-score (r = 0.809, p = 0.001) correlated with the CFQ-Rresp but the FEV₁ z-score (r = 0.249, p = 0.051, figure 2C) did not. The LCI z-score also correlated with the CFCS (r = 0.565, p < 0.001, figure 2B), as did the FEV₁ z-score (r = −0.411, p = 0.002, figure 2D) and FEF₂₅₋₇₅ z-score (r = 0.462, p < 0.001). A separate analysis of children (<12 years) and adolescents (≥12 years) showed similar associations (see online supplementary table E1).

When the analysis was restricted to the subgroup of 53 patients with a normal FEV₁ (z-score > −2), there remained a correlation between the LCI z-score and CFQ-Rresp (r = −0.282, p = 0.043) and between the LCI z-score and the CFCS (r = 0.445, p = 0.002), but not between the FEV₁ z-score and the CFQ-Rresp (r = 0.016, p = 0.908) or CFCS (r = −0.17, p = 0.260).

Pulmonary exacerbations

Of the 63 patients, 26 (41%) received a total of 48 courses of intravenous antibiotics for a PE in the follow-up period after the baseline LCI measurement. Follow-up time was 365 days for 60 patients and 349, 223 and 177 days for the three other patients. The time to the first PE decreased with worsening LCI quartile (p < 0.001, figure 3A) and worsening FEV₁ z-score quartile (p = 0.002, figure 3B) or FEF₂₅₋₇₅ z-score quartile (p = 0.001, see online supplementary figure E3A). The time to the first PE also decreased with worsening CFQ-Rresp quartile (p = 0.001, see online supplementary figure E4A) and with worsening CFCS quartile (p = 0.001, see online supplementary figure E4B).

When the subgroup of patients with a normal FEV₁ was analysed (n = 53), a difference in time to first PE among the LCI quartiles persisted (p = 0.047, figure 3C), although this was not significant.

Table 1 Baseline characteristics of study cohort

<table>
<thead>
<tr>
<th></th>
<th>Patients with CF</th>
<th>Control subjects</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Included</td>
<td>Not included</td>
</tr>
<tr>
<td>N</td>
<td>63</td>
<td>23</td>
</tr>
<tr>
<td>Age in years, median (range)</td>
<td>12.4 (5.3–18.8)</td>
<td>14.0 (4.5–19.6)</td>
</tr>
<tr>
<td>&lt;12 years</td>
<td>31 (49%)</td>
<td>10 (43%)</td>
</tr>
<tr>
<td>Male</td>
<td>33 (52%)</td>
<td>10 (43%)</td>
</tr>
<tr>
<td>F508del homozygote</td>
<td>40 (63%)</td>
<td>12 (52%)</td>
</tr>
<tr>
<td>Pancreatic sufficient</td>
<td>6 (10%)</td>
<td>4 (17%)</td>
</tr>
<tr>
<td>Colonisation status with Pseudomonas aeruginosa†</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chronic</td>
<td>8 (13%)</td>
<td>2 (9%)</td>
</tr>
<tr>
<td>Intermittent</td>
<td>12 (19%)</td>
<td>3 (13%)</td>
</tr>
<tr>
<td>Weight z-score</td>
<td>−0.51 (0.94)</td>
<td>−0.40 (1.17)</td>
</tr>
<tr>
<td>Height z-score</td>
<td>−0.41 (0.98)</td>
<td>−0.73 (1.21)</td>
</tr>
<tr>
<td>BMI z-score</td>
<td>−0.34 (0.90)</td>
<td>0.04 (0.95)</td>
</tr>
<tr>
<td>FEV₁ % predicted</td>
<td>89.4 (15.7)</td>
<td>86.3 (25.7)</td>
</tr>
<tr>
<td>FEV₁ z-score</td>
<td>−0.88 (1.29)</td>
<td>−1.09 (2.10)</td>
</tr>
<tr>
<td>FEF₂₅₋₇₅ % predicted</td>
<td>79.9 (30.5)</td>
<td>74.3 (42.1)</td>
</tr>
<tr>
<td>FEF₂₅₋₇₅ z-score</td>
<td>−1.04 (1.53)</td>
<td>−1.37 (2.08)</td>
</tr>
<tr>
<td>LCI</td>
<td>10.78 (3.09)</td>
<td>7.30 (2.09)</td>
</tr>
<tr>
<td>LCI z-score</td>
<td>6.97 (6.20)</td>
<td>0.00 (1.00)</td>
</tr>
<tr>
<td>CFQ-Rresp median (range)</td>
<td>89 (34–100)</td>
<td>15 (10–30)</td>
</tr>
<tr>
<td>CFCS, median (range)</td>
<td>15 (10–30)</td>
<td></td>
</tr>
</tbody>
</table>

*Data presented as n (%) or mean (SD) unless otherwise specified.
†During the previous 12 months according to the definition of Lee et al.²³

BMI, body mass index; CF, cystic fibrosis; CFCS, Cystic Fibrosis Clinical Score; CFQ-Rresp, respiratory domain of the Cystic Fibrosis Questionnaire-Revised; FEF₂₅₋₇₅, forced expiratory flow between 25% and 75% of forced vital capacity; FEV₁, forced expiratory volume in 1 s; LCI, lung clearance index.
observed for the FEV\textsubscript{1} z-score quartiles (p=0.113, figure 3D) or among the FEF\textsubscript{25−75} z-score quartiles (p=0.452, see online supplementary figure E3B).

The univariate regression analysis (table 2) showed that the annual PE rate was higher among patients in the worst quartile compared with those in the best quartile for LCI (p=0.003). This difference was also observed when comparing the FEV\textsubscript{1} (p=0.016) or FEF\textsubscript{25−75} z-score quartiles (p=0.020). Restricting the analysis to the subgroup of patients with a normal FEV\textsubscript{1}, the patients in the worst LCI quartile still had a higher PE rate than those in the best LCI quartile (p=0.043), whereas the PE rate was not different between patients in the worst FEV\textsubscript{1} z-score quartile (p=0.280) or FEF\textsubscript{25−75} quartile (p=0.453) compared with those in the best quartiles.

Age, isolation of \textit{P aeruginosa} in the previous year, LCI z-score, FEV\textsubscript{1} z-score, FEF\textsubscript{25−75} z-score, CFCS and CFQ-R\textsubscript{resp} were used as predictors for the multivariate prediction model because they were found to be significant predictors of the PE rate in the univariate analysis (table 3). The backward stepwise approach identified the LCI z-score (p=0.001) as the only independent predictor of the PE rate in the year following the baseline assessment. An increase in the LCI z-score of 1 resulted in an increase in the PE rate by 12.0% (95% CI 5.0% to 19.5%). Restricting the analysis to patients with a normal FEV\textsubscript{1}, the LCI z-score remained a predictor of the PE rate in the following year (rate ratio (RR) 1.12, 95% CI 1.02 to 1.23, p=0.017) and a trend was shown for the FEV\textsubscript{1} z-score (RR 0.57, 95% CI 0.31 to 1.07).
and the FEF$_{25-75}$ z-score (RR 0.72, 95% CI 0.47 to 1.11, $p=0.134$).

**DISCUSSION**

In the present study we have demonstrated a clear association between the baseline LCI and subsequent PEs in a cohort of 5–19-year-old patients with CF. We also documented a correlation between the LCI and the CFQ-Rresp, a validated patient-reported outcome. This result demonstrates the association between the LCI and clinically meaningful endpoints, which is a necessary step in the validation of LCI as a surrogate endpoint for clinical trials involving patients with CF. In the subgroup of patients with a normal FEV$_1$, the LCI was more closely related to PEs and CFQ-R$_{resp}$ than were FEV$_1$ and FEF$_{25-75}$, highlighting the added value of the LCI over FEV$_1$ and FEF$_{25-75}$ in the group of patients with early CF lung disease.

**LCI and PEs**

The cohort included in this study is representative of the current spectrum of lung disease in children and young adults.
lung disease. At the severe end of the spectrum of lung function abnormalities, PEs were predicted by FEV\textsubscript{1} as all patients with an abnormal FEV\textsubscript{1} had at least one PE in the 1-year follow-up period. This observation was not unexpected because LCI remained as the single predictor of PEs in patients with a normal FEV\textsubscript{1}. Only two of 11 patients with a normal LCI were treated with intravenous antibiotics for a PE: one patient was a 10-year-old boy with an expected superior sensitivity of the LCI to detect early disease spectrum.

With the expected superior sensitivity of the LCI to detect early lung disease, \textsuperscript{8} at the severe end of the spectrum of lung function abnormalities, PEs were predicted by FEV\textsubscript{1} as all patients with an abnormal FEV\textsubscript{1} had at least one PE in the 1-year follow-up period. This observation was not unexpected because LCI remained as the single predictor of PEs in patients with a normal FEV\textsubscript{1}. Only two of 11 patients with a normal LCI were treated with intravenous antibiotics for a PE: one patient was a 10-year-old boy with an expected superior sensitivity of the LCI to detect early lung disease. \textsuperscript{8} At the severe end of the spectrum of lung function abnormalities, PEs were predicted by FEV\textsubscript{1} as all patients with an abnormal FEV\textsubscript{1} had at least one PE in the 1-year follow-up period. This observation was not unexpected because LCI remained as the single predictor of PEs in patients with a normal FEV\textsubscript{1}. Only two of 11 patients with a normal LCI were treated with intravenous antibiotics for a PE: one patient was a 10-year-old boy with an expected superior sensitivity of the LCI to detect early lung disease. \textsuperscript{8}

As there is no universally accepted definition for PE, \textsuperscript{25} we chose to use a conservative definition including only episodes treated with intravenous antibiotics. PEs treated with intravenous antibiotics are known to have an impact on long-term outcomes \textsuperscript{26} and patient-reported outcomes, \textsuperscript{27} which suggests that our definition of PE represented a relevant clinical outcome. Episodes of treatment with oral antibiotics were not considered in the present study as their impact on outcomes has not been established and because oral antibiotics are also prescribed for positive airway cultures, regardless of clinical symptoms. The decision to administer intravenous antibiotics was made during multidisciplinary team meetings in a single centre, thereby mimicking the differences in practice among physicians. Using this definition, 40% of our patients experienced a PE, which is in line with previously published data in similar age cohorts. \textsuperscript{24}

### Table 2: Annual rate of PEs and rate ratios (RRs) from the univariate negative binomial regression models by quartiles of LCI and FEV\textsubscript{1} z-scores in all patients and in patients with a normal FEV\textsubscript{1}

| Parameter | All patients (n=63) | | Patients with normal FEV\textsubscript{1} (n=53) | |
|-----------|---------------------|---------------------|
|          | Annual PE rate (95% CI) | p Value* | RR (95% CI) | p Value* | LCI z-score | |
| ≤−2.4    | 0.1 (0.0–0.6) | 1 | | 0.1 (0.0–0.6) | 1 |
| 2.6–5.0  | 0.4 (0.1–1.0) | 0.088 | 2.9 (0.5–16.5) | 0.238 | 2.4–3.9 | 0.5 (0.2–1.2) | 0.462 | 3.3 (0.6–19.04) | 0.186 |
| 5.3–10.6 | 0.7 (0.3–1.6) | 0.016 | 5.4 (1.0–29.0) | 0.045 | 4.2–6.8 | 0.6 (0.2–1.5) | 0.302 | 4.1 (0.7–23.4) | 0.114 |
| ≥11.4    | 1.8 (1.0–3.3) | 0.003 | 13.6 (2.8–67.1) | 0.001 | ≥7.1 | 0.9 (0.4–2.0) | 0.043 | 6.5 (1.2–34.5) | 0.029 |

### Table 3: Univariate negative binomial regression models for pulmonary exacerbations in the year following the baseline measurement

<table>
<thead>
<tr>
<th>Predictor</th>
<th>β</th>
<th>SE (β)</th>
<th>Wald χ\textsuperscript{2}</th>
<th>p Value</th>
<th>Rate ratio</th>
<th>95% CI of rate ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>LCI z-score</td>
<td>0.11</td>
<td>0.03</td>
<td>11.9</td>
<td>0.001</td>
<td>1.12</td>
<td>1.05–1.19</td>
</tr>
<tr>
<td>FEV\textsubscript{1} z-score</td>
<td>−0.54</td>
<td>0.17</td>
<td>10.3</td>
<td>0.001</td>
<td>0.58</td>
<td>0.42–0.81</td>
</tr>
<tr>
<td>CFCS</td>
<td>0.15</td>
<td>0.05</td>
<td>8.3</td>
<td>0.004</td>
<td>1.16</td>
<td>1.05–1.29</td>
</tr>
<tr>
<td>FEF\textsubscript{25–75} z-score</td>
<td>−0.38</td>
<td>0.13</td>
<td>8.3</td>
<td>0.004</td>
<td>0.69</td>
<td>0.53–0.89</td>
</tr>
<tr>
<td>Ps.a positive*</td>
<td>0.90</td>
<td>0.40</td>
<td>5.5</td>
<td>0.023</td>
<td>2.46</td>
<td>1.13–5.36</td>
</tr>
<tr>
<td>CFQ-Rresp</td>
<td>−0.03</td>
<td>0.01</td>
<td>5.3</td>
<td>0.021</td>
<td>0.97</td>
<td>0.94–0.99</td>
</tr>
<tr>
<td>Age in years</td>
<td>0.10</td>
<td>0.05</td>
<td>3.9</td>
<td>0.049</td>
<td>1.10</td>
<td>1.00–1.22</td>
</tr>
<tr>
<td>Female gender</td>
<td>0.50</td>
<td>0.39</td>
<td>1.6</td>
<td>0.199</td>
<td>1.65</td>
<td>0.77–3.53</td>
</tr>
</tbody>
</table>

* p Value for the difference among each quartile and the best quartile from the univariate negative binomial regression models.

CF, cystic fibrosis; FEV\textsubscript{1}, forced expiratory volume in 1 s; LCI, lung clearance index; PE, pulmonary exacerbation.

Patient-reported outcomes

Correlations with patient-reported outcomes have been reported for FEV\textsubscript{1}, \textsuperscript{28} total CT score, \textsuperscript{29} and bronchiectasis and air trapping scores. \textsuperscript{30} Data on the correlations among LCI, symptoms and patient-reported outcomes are sparse. Bakker et al\textsuperscript{31} measured the LCI and cough counts over two nights in preschool children with CF and found no correlation between these variables. Aurora et al\textsuperscript{15} found a correlation between the LCI and recent cough in preschool children with CF. However, the study by Amin et al\textsuperscript{12} found no correlation between the LCI and the CFQ-R\textsubscript{resp} in a small group of patients. We showed that the LCI correlated with the CFQ-R\textsubscript{resp} even in the group with a normal FEV\textsubscript{1} even though FEV\textsubscript{1} did not, which again highlights the superiority of LCI over FEV\textsubscript{1} at the milder end of the CF disease spectrum.

Whether other spirometric indices such as the maximum expiratory flow at 50% of FVC, \textsuperscript{32} forced expiratory flow at 75% of FVC, \textsuperscript{29} and FEF\textsubscript{25–75} \textsuperscript{12} could be more sensitive for early CF lung disease than FEV\textsubscript{1} has been debated. In the present cohort only a minority of patients (6 of 53) with a normal FEV\textsubscript{1} had an
abnormal FEV₁, suggesting that FEF₂₅₋₇₅ adds little information to that provided by FEV₁. In addition, FEF₂₅₋₇₅ correlated with CFQ-RRESP, and was predictive of PE but, unlike LCI, was not a predictor of PE in patients with a normal FEV₁.

**Future developments**

Demonstrating a relationship with subsequent mortality would validate the LCI as a surrogate outcome measure. However, whether initial disease progression as shown by changes in the LCI at a younger age will continue during adulthood and lead to an untimely death remains unclear although, in adults, the LCI has been correlated with FEV₁, a strong predictor of mortality. After establishment of the prognostic value of LCI for occurrence of PEs, more information is needed about fluctuations of LCI in patients with CF, the effect of exacerbations on LCI and the timing and magnitude of LCI improvement with interventions to determine what represents a clinically meaningful change.

This study was not designed to explore whether or how LCI can be used to guide therapeutic decisions in an individual patient. It is not known how much decline would indicate the need for additional treatment or how much increase would constitute a significant improvement.

The placement of the LCI in the outcome toolbox requires further refinement, and it needs to be evaluated how this measure compares with other surrogate outcomes such as the chest CT scan score. However, the LCI appears to fit well as an endpoint in children with CF and mild lung disease in whom FEV₁ lacks sensitivity and in whom the need for repeated measurements contraindicates chest CT due to radiation risks.

In conclusion, the present study shows that the LCI is predictive of PE and is related to patient-reported outcomes and symptoms in 5–19-year-old patients with CF, even in the subgroup with a normal FEV₁. These results add to the validation of the LCI as a surrogate endpoint for use in clinical trials.

**Acknowledgements**

The authors thank the children and families participating in the study. They also thank Kris Colpaert, Linda Boulanger, Nathalie Feytaerts and Jill Ophoff for technical assistance and help with data collection and management.

**Contributors**

Acquisition of data: FV, MP, MB and KDB. Analysis and interpretation of data: FV and KDB. Drafting or revising the paper for important intellectual content and final approval of the paper: FV, MP, MB, TH and KDB.

**Funding**

This study was supported by a grant from the ‘Belgische Vereniging voor de Strijd tegen Mucoviscidose’ and ‘Klinisch Onderzoeksfonds’.

**Competing interests**

None.

**Ethics approval**

The study was approved by the ethics committee of the University of Leuven.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

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Lung clearance index predicts pulmonary exacerbations in young patients with cystic fibrosis

François Vermeulen, Marijke Proesmans, Mieke Boon, Trudy Havermans and Kris De Boeck

Thorax 2014 69: 39-45 originally published online September 10, 2013
doi: 10.1136/thoraxjnl-2013-203807

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