Critical evaluation of computerised x ray planimetry for the measurement of lung volumes

D P S Spence, Y J Kelly, J Ahmed, P M A Calverley, M G Pearson

Abstract

Background – Computerised x ray planimetry has been advocated as an alternative to body plethysmography and helium dilution for measuring static lung volumes. The accuracy and reproducibility of this method has been assessed in comparison with these standard methods.

Methods – Plethysmographic and planimetric measurements of total lung capacity (TLC) and functional residual capacity (FRC) were made in 10 normal subjects and in 12 patients with chronic obstructive pulmonary disease (COPD), with additional helium dilution measurements in the latter 12 patients.

Results – Mean lung volumes (TLC and FRC) for groups of subjects measured by planimetry and by plethysmography were similar in both groups and larger than the helium dilution measurement in patients with COPD. Intraindividual agreement between planimetry and plethysmography was poor, however, with a wide confidence interval (–2.2 to +2.3 l). The planimetry did not measure reliably changes in volume from TLC to FRC in individuals.

Conclusions – Mean lung volumes measured by planimetry in a group of patients probably reflect a regression to the mean of the computer algorithm rather than accurate TLC estimation. The technique is not yet robust enough to replace the established techniques of helium dilution or plethysmography.

(Keywords: x ray planimetry, plethysmography, helium dilution, lung volumes.)

Various techniques of increasing sophistication have been described to derive lung volumes from the chest radiograph.1–7 The most recent planimetric method8 defines the boundaries of the chest wall, heart, diaphragm, and spine on posteroanterior and lateral radiographs and has been adapted into a commercially available computerised package (PK Morgan Ltd, Rainham, Kent, UK). The radiographic method has the theoretical advantage in patients with chronic obstructive pulmonary disease (COPD) of overcoming the artefactual errors caused by prolonged time constants for gas mixing and pressure equilibration that can confound both helium dilution and plethysmographic techniques.8–11 Radiographic lung volume estimation could save a visit to the respiratory laboratory, is inexpensive, and has been used in recently published epidemiological studies.12 Such practical advantages are only of value, however, if the radiographic measurements are both reliable and well validated against more established techniques. Limited data are available on earlier radiographic techniques but none concerning the computerised planimeter.

We have evaluated the planimetric technique in both normal individuals and in patients with COPD and have examined (1) whether the planimetric lung volume measurements can be measured reproducibly by different observers; (2) if radiographic lung volumes are comparable with the helium dilution and plethysmographic values; (3) whether the radiographic method can accurately detect changes in lung volume; and (4) if the posture adopted for the chest radiograph affects the results.

Methods

Patients

Ten healthy male volunteers (mean age 33 (3) years) were recruited from the hospital staff and 12 patients (nine men) with a diagnosis of COPD13 were recruited from the outpatient clinic (table 1). Written informed consent was obtained from all subjects and the study was approved by the district ethical committee.

Protocol

Posteroanterior and lateral chest radiographs were taken with the subject seated to be comparable with the plethysmographic and helium dilution methods. Ventilation was monitored during the procedure by having the subject rebreathe from a water sealed spirometer (PK Morgan Ltd) incorporating a CO2 absorber. Oxygen was added to the system to maintain a constant volume of gas within the apparatus.

Table 1 Mean (SD) demographic data on all subjects

<table>
<thead>
<tr>
<th></th>
<th>Normal volunteers</th>
<th>COPD patients</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(n = 10)</td>
<td>(n = 12)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>32 (2.7)</td>
<td>61 (3.3)</td>
</tr>
<tr>
<td>FEV1 (L)</td>
<td>4 (0.4)</td>
<td>0.9 (0.07)</td>
</tr>
<tr>
<td>FEV1 (% predicted)</td>
<td>102 (5)</td>
<td>28 (8)</td>
</tr>
<tr>
<td>FRC (L)*</td>
<td>3.6 (0.85)</td>
<td>6.33 (0.29)</td>
</tr>
<tr>
<td>TLC (L)†</td>
<td>7.08 (1.12)</td>
<td>7.68 (1.24)</td>
</tr>
</tbody>
</table>

FEV1 = forced expiratory volume in one second; FRC = functional residual capacity; TLC = total lung capacity; COPD = chronic obstructive lung disease.

* Measured by body plethysmography.
Respiration was monitored by watching the water sealed spirometer. Subjects were instructed to breathe in fully until a plateau was observed on the spirometric trace, at which point the posteroanterior radiograph at total lung capacity (TLC) was taken. Subjects were then told to relax and breathe normally and, after a few breaths, the functional residual capacity (FRC) posteroanterior radiograph was taken, the breath being held at end tidal lung volume as judged from the spirometer. The procedure was repeated to obtain a pair of lateral chest radiographs. The distance from x-ray tube to film was six feet throughout.

**PLANIMETRIC LUNG VOLUMES**

In the normal volunteers planimetric x-ray lung volumes were determined by a single observer (DS). In the patients with COPD lung volumes were independently determined from the same pairs of radiographs by DS and a second observer (YJK).

Briefly, the measurement procedure comprises tracing the pleural outline and those of the other intrathoracic structures (fig 1). The posteroanterior and lateral radiographs are aligned using the arch of the aorta as a common reference point. The computer program then divides the resulting shapes into 200 slices, calculates the volume of each slice and, thus, the total chest volume. From this the volumes of the heart, great vessels, subdiaphragmatic region, and spine (including paravertebral musculature) are subtracted to leave the total lung volume. A correction factor is applied to this total lung volume to take account of solid tissue and volume of blood within the lungs. The net result is the calculated volume of air within the lungs.

**CONVENTIONAL LUNG VOLUME MEASUREMENTS**

Lung volumes were measured in a constant volume body plethysmograph (PK Morgan Ltd) with the subject panting at 1 Hz. In both the normal volunteers and patients measurements were made whilst supporting the cheeks with the hands, and in the normal subjects additional measurements were made with the arms raised to mimic the position adopted for the lateral radiograph. In the patients lung volumes were also measured by the standard rebreathing helium dilution technique.

**STATISTICAL ANALYSIS**

Data are expressed as mean (SD) unless otherwise stated. Comparisons between measurement techniques were evaluated using the methods recommended by Bland and Altman with 95% confidence intervals (CI). For comparison with previously published work on x-ray planimetry these results were compared with simple linear regression analysis.

**REPRODUCIBILITY OF CONVENTIONAL MEASUREMENTS**

The reproducibility of body plethysmography and of helium dilution lung volumes in our

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**Table 2** Mean (SD) values for TLC and FRC in normal subjects and patients with COPD, and 95% CI of agreement between different measures

<table>
<thead>
<tr>
<th></th>
<th>Mean TLC (l)</th>
<th>Mean FRC (l)</th>
<th>95% CI for an individual value</th>
<th>95% CI for an individual value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Normal subjects</strong> (n=10)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>x ray planimetry</td>
<td>7·13 (1·02)</td>
<td>4·62 (0·85)</td>
<td>−2·19 to +2·29</td>
<td>−0·06 to +2·10</td>
</tr>
<tr>
<td>Body plethysmography (seated)</td>
<td>7·08 (1·12)</td>
<td>3·60 (1·01)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Body plethysmography (arms raised)</td>
<td>7·32 (0·97)</td>
<td>4·01 (0·87)</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>COPD subjects</strong> (n=12)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>x ray planimetry</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Observer 1</td>
<td>7·24 (1·46)</td>
<td>6·17 (1·63)</td>
<td>−0·67 to +0·67</td>
<td>−0·39 to +0·57</td>
</tr>
<tr>
<td>Observer 2</td>
<td>7·23 (1·46)</td>
<td>6·27 (1·59)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Body plethysmography (seated)</td>
<td>7·68 (1·24)</td>
<td>6·33 (0·99)</td>
<td>−2·40 to +1·52</td>
<td>−2·42 to +2·12</td>
</tr>
<tr>
<td>Helium dilution</td>
<td>6·60 (1·10)</td>
<td>4·95 (0·86)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

TLC = total lung capacity; FRC = functional residual capacity; COPD = chronic obstructive pulmonary disease.
laboratory have been assessed in a separate study of 32 patients with COPD of similar severity (age 62.5 (7-5) years, FEV1 0.77 (0.24) l), each studied on four occasions. The coefficient of variation for both techniques was less than 5%, yielding a confidence interval for an individual measurement by body plethysmography of 0.31 to 0.421.

Results

NORMAL VOLUNTEERS (table 2)
Mean TLC values measured by plethysmography and by x ray planimetry were similar for the whole group. However, the Bland and Altman approach showed that there was poor concordance within individual subjects such that the differences had a 95% CI of -2.19 to +2.29 l (fig 2). Mean FRC measured with the planimeter was higher than with the plethysmograph, but again the 95% CI of differences within individuals was wide (-0.06 to +2.71).

The mean inspiratory capacity from FRC to TLC measured on the spirometer was similar to that derived from differences in radiographs at FRC and TLC (2.88 (0.34) l), but again there were considerable discrepancies within individuals (95% CI of difference in techniques -1.75 to +2.49 l).

A

Difference between planimetric and plethysmographic measurements (l)

B

95% CI

95% CI

Discussion
All biological measurements have an inherent variability and, to interpret a result in an individual patient, the degree of confidence which can be placed on a given value must be known. Measurements of lung volume by helium dilution and by body plethysmography have been used for many years and have a coefficient of variation of between 5% and 10%. In our laboratory we have found a coefficient of variation for repeated measurements on separate days in patients with COPD of less than 5% for both techniques (95% CI ±0.6 l). Moreover, both helium dilution and plethysmographic measurements are supported by an extensive amount of literature which shows that the data they yield are clinically reproducible and able to detect reliably the changes in lung function resulting from disease or treatment.

We have shown that the mean lung volume for two groups of subjects, with and without COPD, are similar whether measured by plethysmography, helium dilution, or by x ray planimetry. In previous studies the individual data have been plotted on an x-y plot and a linear regression performed to assess reproducibility. Our data treated in this way yield a linear regression correlation coefficient (r) of 0.95 which is very similar to previous work. It is therefore tempting to argue that the method has been validated since mean values agree and individual values are significantly correlated.

However, the statistical validity of this approach has been challenged as it does not compare the repeatability of the measurement alone, but reflects other factors such as body size. Tall people tend to have larger lung volumes than small people no matter what technique is used. If the subject group includes both tall and short people then a simple x-y plot and linear regression will inevitably appear to correlate well because of the relationship of...
volume to height. Intertest differences therefore cannot be assessed by linear regression. Bland and Altman showed how the intrapersonal differences between measurement techniques should be assessed and, when applied to our data, this method shows a very wide 95% CI for individuals. Thus, for TLC it indicates that, if the first value is known, the second technique had a 95% chance of being within 2-21 of the first.

The internal reproducibility of the x ray planimetry measurement as recorded by two different observers on the same pair of radiographs was good, suggesting that the mechanical aspects of the planimeter and translation of radiograph to computer is not a problem.

Plethysmographic values are known to be consistently larger than helium dilution values for good physiological reasons and, not surprisingly, our results in patients with COPD confirmed this.

The concordance of mean values between x ray planimetry and other techniques probably reflects the fact that the algorithm is showing a regression to the mean that hides individual variation. For individuals there are a number of possible explanations for the poor concordance with other techniques. The radiographic exposure will vary with the radiographer, and a different grey scale will result in different lung outlines being defined. Thus, planimetry volumes measured by two observers from the same radiograph agree more closely than does planimetry with other techniques. Secondly, the computer algorithm makes a number of assumptions about the geometry of the chest. It cannot correct for individuals of different shape or for the change in shape that accompanies either a change in volume or change due to disease. Thirdly, the computer assumes that a patient has breathed in to TLC when the radiograph was taken. Some patients, for varying reasons, find it difficult to comply with simple commands and the radiographer has no method of checking that full inspiration has occurred.

The radiographic technique algorithms were derived to measure TLC. It is important for a measurement technique to be able to record changes in volume that occur as physiology and anatomy alter with disease states. The planimeter was not sufficiently sensitive to record reliably changes in volume between TLC and FRC compared with a spirometer. The wide confidence interval for agreement within an individual must throw doubt on the ability of planimetry to measure accurately or reliably change due to disease. Previous studies used less sophisticated methods of calculating volume from radiographs, relying on as few as five measurements from the paired films. It is likely that these simple systems will be even more prone to error.

Lastly, with the use of the body plethysmograph, we showed that raising the arm position to mimic the position for a lateral chest radiograph results in a small but significant increase of 0-251 in TLC which means that posteroanterior and lateral radiographs are actually being measured at different values of TLC, which is a further complication for a computer algorithm.

In conclusion, radiographic estimation of lung volume may have theoretical advantages, especially in patients with airflow limitation, but when the method is critically assessed with the appropriate statistical technique the values in individuals are not comparable to established techniques. The variability within an individual is too large to be clinically useful. Moreover, the inability to detect accurately change in volume within the same person suggests that the radiographic measurement may not be a reliable indicator of either worsening disease or of a response to treatment. Even its value in following changes in group mean data cannot be assumed until it is compared with standard techniques in the same population of individuals in months or years. Until the method has been shown to reflect reliably within-patient changes resulting from disease or treatment, there must be some doubt as to whether it is valid for use as part of epidemiological surveys. The planimeter itself is easy to use and internally reproducible, but the computer algorithm needs further refinement to make it clinically relevant.
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