Influence of emphysema distribution on pulmonary function parameters in COPD patients

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ABSTRACT

Objective: To evaluate the impact that the distribution of emphysema has on clinical and functional severity in patients with COPD. Methods: The distribution of the emphysema was analyzed in COPD patients, who were classified according to a 5-point visual classification system of lung CT findings. We assessed the influence of emphysema distribution type on the clinical and functional presentation of COPD. We also evaluated hypoxemia after the six-minute walk test (6MWVT) and determined the six-minute walk distance (6MWD).

Results: Eighty-six patients were included. The mean age was 65.2 ± 12.2 years, 91.9% were male, and all but one were smokers (mean smoking history, 62.7 ± 38.4 pack-years). The emphysema distribution was categorized as obviously upper lung-predominant (type 1), in 36.0% of the patients; slightly upper lung-predominant (type 2), in 25.6%; homogeneous between the upper and lower lung (type 3), in 16.3%; and slightly lower lung-predominant (type 4), in 22.1%. Type 2 emphysema distribution was associated with lower FEV₁, FVC, FEV₁/FVC ratio, and DLCO. In comparison with the type 1 patients, the type 4 patients were more likely to have an FEV₁ < 85% of the predicted value (OR = 6.91, 95% CI: 1.43-32.64; p = 0.016), a 6MWD < 350 m (OR = 6.36, 95% CI: 1.26-32.18; p = 0.025), and post-6MWVT hypoxemia (OR = 32.66, 95% CI: 3.26-326.84; p = 0.003). The type 3 patients had a higher RV/TLC ratio, although the difference was not significant.

Conclusions: The severity of COPD appears to be greater in type 4 patients, and type 3 patients tend to have greater hyperinflation. The distribution of emphysema could have a major impact on functional parameters and should be considered in the evaluation of COPD patients.

Keywords: Pulmonary disease, chronic obstructive; Pulmonary emphysema; Respiratory function tests; Tomography, X-ray computed.

INTRODUCTION

The lung disease known as COPD is characterized by persistent airflow limitation that is usually progressive, consisting of a combination of small airways disease (obstructive bronchiolitis) and parenchymal destruction (emphysema).¹ There is increasing evidence to suggest that distinguishing different phenotypic profiles of patients with COPD has prognostic and therapeutic implications.²⁻⁴ In fact, COPD patients with confirmed emphysema have more severe lung function impairment, more intense airway inflammation, and possibly more important extrapulmonary disability than do those without emphysema.²⁻⁴,⁶ The lung hyperinflation caused by the loss of lung elastic recoil has been associated with limitations in the functional capacity of these patients.⁷⁻⁸ In addition, the destruction of the alveolar-capillary membrane in emphysema is responsible for more profound hypoxemia.⁹⁻¹₀

Advances in CT scanning and image processing software have allowed the precise measurement of the extent of low-attenuation areas corresponding to emphysema. In validation studies, the results obtained with these techniques have been found to correlate well with pathologic and functional features.¹⁰⁻¹³ This kind of assessment has been mainly used in order to evaluate patients for lung volume reduction procedures and to monitor replacement therapy in alpha-1 antitrypsin-deficient patients.¹⁴⁻¹⁶ However, quantifying emphysema might have broader utility, given that some reports have shown that the heterogeneity of the distribution of parenchymal damage might be associated with different degrees of clinical severity.¹³⁻¹⁷⁻¹⁹ Nevertheless, the results are contradictory, which might be attributable to the different methods that have been used in those analyses. The majority of authors have employed computer-assisted measurements, which are expensive and not widely available. In order to promote a definitive widespread use of imaging data in the clinical evaluation of patients with emphysema, we believe that there is also a need to standardize qualitative methods.

The aim of the present study was to evaluate the impact that the distribution of emphysema has on clinical and functional features in COPD patients. In order to test our hypothesis, we used a visual classification system to categorize patients according to the regional distribution of their emphysema.

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**METHODS**

**Study subjects**

This was a cross-sectional observational study involving COPD patients with emphysema, recruited between August of 2011 and August of 2012 from the pulmonology outpatient clinic of the **Centro Hospitalar de São João**, a tertiary care medical center located in the city of Porto, Portugal. We included patients with pulmonary emphysema and any degree of airflow limitation who had been clinically stable in the 3 months prior to their inclusion in the study. The inclusion criteria were having a post-bronchodilator FEV1/FVC < 0.70 and showing evidence of emphysema on visual inspection of CT images, estimated to involve > 25% of the lung parenchyma. Patients with a history of asthma, bronchiectasis, tuberculosis sequelae, lung fibrosis, thoracic surgery, or other confounding diseases were excluded (Figure 1). The study was approved by the local research ethics committee, and all patients gave written informed consent.

**Clinical and pulmonary function assessment**

We recorded demographic and anthropometric data, namely age, gender, and BMI. Patients also underwent clinical evaluation, which included the completion of the COPD Assessment Test (CAT) and the modified Medical Research Council (mMRC) scale (for the determination of dyspnea severity), as well as the evaluation of smoking status (current smoker, former smoker, or nonsmoker), smoking history (in pack-years), the presence of significant comorbidities, and current medication use. The number of COPD exacerbations in the last year was retrospectively obtained by patient recall, and, in most cases, hospital records were used in order to corroborate the information. Each patient was submitted to spirometry (MasterScreen™ Body; Jaeger, Würzburg, Germany), lung volumes and DLCO was submitted to spirometry (MasterScreen™ Body; Jaeger, Würzburg, Germany), lung volumes and DLCO were performed using the methodology described by the American Thoracic Society. Arterial blood gases were measured (RapidLab™ 1265; Siemens, Munich, Germany) after a minimum 30-min rest period in a sitting position. We defined hypoxemia as a PaO2 < 60 mmHg at an FiO2 of 0.21.

**CT evaluation**

All patients underwent multidetector CT of the chest at suspended full inspiration, from the thoracic inlet to the adrenal glands, using a 64-detector row scanner (Somatom Sensation 64; Siemens Healthcare, Erlangen, Germany). The following imaging parameters were used: tube voltage, 120 kVp; tube current, 40 mAs; rotation time, 0.33 s; pitch, 1.3; detector collimation, 32 × 0.6 mm; and slice acquisition by means of a z-flying focal spot, 64 × 0.6 mm. No contrast media were used. From the raw data, 1 mm-thick sections were obtained using a soft tissue kernel reconstruction (B50f; Siemens Healthcare). For the subjects submitted to multiple CT scans, the one performed the closest to study enrollment was used.

Two thoracic radiologists independently reviewed the CT imaging studies. Both were blinded to the clinical information of the patients. Disagreement between the two radiologists was resolved by consensus. They reviewed CT images on the coronal and sagittal planes to assess the heterogeneity of emphysematous changes in an apical-to-caudal direction. For image interpretation, we used a window level of −700 to −900 HU and a window width of 600-1,600 HU. A five-point visual classification system was applied as previously described. This qualitative evaluation ranks pulmonary emphysema according to its predominant distribution, as follows: type 1, obviously predominant in the upper lung; type 2, somewhat predominant in the upper lung; type 3, equal extent in the upper and lower lung (homogeneous distribution); type 4, somewhat predominant in the lower lung; and type 5, obviously predominant in the lower lung.

**Statistical analysis**

Variables with normal distribution are expressed as means and standard deviations, whereas those with non-normal distribution are expressed as median and interquartile range (25th to 75th percentile) and categorical variables are expressed as absolute values and proportion. The Student’s t-test for independent samples was used in order to compare variables with normal distribution, and the Mann-Whitney U rank test was used in order to compare variables with non-normal distribution. The Pearson’s chi-square test was used for categorical variables. One-way ANOVA was used in order to compare the emphysema distribution groups, together with Tukey’s post hoc test to identify significant differences. Odds ratios and the corresponding 95% confidence intervals were calculated using binary logistic regression. Odds ratios were adjusted for age and BMI. Statistical significance was set at p < 0.05 (two-tailed), and all statistical analyses were performed with the SPSS Statistics software package, version 19.0 (IBM Corporation, Armonk, NY, USA).

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**Figure 1.** Flowchart for the selection of the participating patients.
RESULTS

During the study period, 86 COPD patients with pulmonary emphysema were selected, the characteristics of whom are shown in Table 1. Male gender was predominant in this population, all but one of the patients were current or former smokers, and 1 patient presented with homozygous PiZZ alpha-1 antitrypsin deficiency. The CAT scores indicated severe symptoms more often than did the mMRC dyspnea scale scores: 56.1% of the patients had CAT scores ≥ 10, whereas only 40.8% had mMRC scale scores ≥ 2. Only 24.4% of the patients had frequent exacerbations (≥ 2 exacerbations in the last year). Hypoxemia was present in 15 patients (18.8%). The characteristics of the patients demonstrated a wide range of airflow limitation, with an even distribution across the Global Initiative for Chronic Obstructive Lung Disease severity classification, which is based on FEV₁—mild, in 27.9%; moderate, in 22.1%; severe, in 29.1%; and very severe, in 20.9%—reflecting the spectrum of the disease encountered in clinical practice. However, there was a clear tendency for hyperinflation to be seen in this group of patients, with a median residual volume/total lung capacity (RV/TLC) ratio of 55.5% and a significant median oxygen desaturation during the 6MWT of 6%.

In most (36.0%) of the patients, the emphysema was obviously predominant in the upper lung (type 1 distribution). The next most common distributions were types 2 and 4 (somewhat predominant in the upper lung and somewhat predominant in the lower lung, seen in 25.6% and 22.1% of the patients, respectively). Type 3 emphysema distribution (homogeneous distribution between the upper and lower lung) was the least common, seen in only 16.3%. None of the patients in our sample were classified as presenting with type 5 emphysema distribution (obviously predominant in the lower lung). The interobserver correlation for emphysema classification scores was good (rₛ = 0.621, p < 0.001).

Figure 2 shows the differences found in the clinical parameters according to CT scan classification of emphysema distribution. The six-minute walk distance (6MWD), post-6MWT oxygen desaturation, FVC, FEV₁, and FEV₁/FVC ratio (in % of the predicted values), as well as DLCO, were found to differ significantly among the groups. Tukey’s post hoc test revealed that there were significant differences in all of the abovementioned variables between the patients classified as type 1 than those classified as type 4. In fact, all of those variables appear to get worse in upper-to-lower predominance direction. Patients classified as type 3 showed the highest RV/TLC ratio, although it did not reach statistical significance (p = 0.064).

The logistic regression analysis for different dimensions of the functional status revealed that type 4 patients had a significantly higher risk for having

| Table 1. Demographic, clinical, and imaging characteristics of selected patients with emphysema-predominant COPD.* |
|-----------------------------------------------|
| Characteristic | (n = 86) |
| Age, years | 65.2 ± 12.2 |
| Gender | 79 (91.9) |
| Male | 7 (8.1) |
| Female | 23.1 ± 4.5 |
| BMI, kg/m² | 54 (38-79) |
| Smoking history, pack-years | 1 (0.5-3.0) |
| mMRC dyspnea scale score | 35 (40.8) |
| mMRC dyspnea scale score ≥ 2 | 12 (7.0-22.5) |
| CAT score | 48 (56.1) |
| ≥ 2 exacerbations in the last year | 21 (24.4) |
| Hypoxemia | 15 (18.8) |
| Post-6MWT desaturation, % | 6 (4.0-9.8) |
| 6MWD, m | 400 (256.3-463.8) |
| FVC, % of predicted | 86.1 ± 24.8 |
| FEV₁ | 50.0 (32.0-83.3) |
| FEV₁/FVC ratio | 45.9 (35.0-63.2) |
| RV | 162.0 (125.1-225.0) |
| TLC | 118.3 ± 24.4 |
| RV/TLC ratio | 55.5 (43.9-67.1) |
| DLCO | 59.0 (40.0-77.7) |
| Emphysema distribution | 31 (36.0) |
| Type 1 (obvious upper-lung predominance) | 22 (25.6) |
| Type 2 (slight upper-lung predominance) | 14 (16.3) |
| Type 3 (equal upper- and lower-lung extent) | 19 (22.1) |

mMRC: modified Medical Research Council; CAT: COPD Assessment Test; 6MWT: six-minute walk test; and 6MWD: six-minute walk distance. *Values are presented as mean ± SD, n (%), or median (interquartile range).
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FEV<sub>1</sub> < 65% of the predicted value (adjusted OR = 6.92; 95% CI: 1.43-33.45; p = 0.016), 6MWD < 350 m (adjusted OR = 6.36; 95% CI: 1.26-32.18; p = 0.025), and hypoxemia (adjusted OR = 32.66; 95% CI: 3.26-326.84; p = 0.003; Table 2). However, none of the different types of emphysema distribution were found to be significant predictors of BMI ≤ 21 kg/m<sup>2</sup>, ≥ 2 exacerbations in the last year, mMRC dyspnea scale score ≥ 2, or post-6MWT oxygen desaturation ≥ 4%.

**DISCUSSION**

Although COPD is a highly heterogeneous disease, its phenotyping can be more precise when CT of the lung parenchyma is combined with an evaluation of the clinical and physiological characteristics. Here, we describe the role of using a qualitative analysis of CT findings in order to determine the distribution of pulmonary emphysema and the potential contribution

*Figure 2. Lung functional characteristics of patients with different emphysema distribution. *Indicates a p value < 0.05. 6MWT: six-minute walk test; type 1: obviously predominant emphysema in upper lung; type 2: somewhat predominant emphysema in upper lung; type 3: equal extent of emphysema in upper and lower lung; and type 4: somewhat predominant emphysema in lower lung.*

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of that distribution to further characterizing the clinical severity of these patients.

Patients with COPD were classified according to a subjective heterogeneity analysis of upper versus lower lung distribution of pulmonary emphysema, using a visual scoring system first described by Chae et al.\(^\text{(18)}\) In their assessments of the regional heterogeneity of the distribution of emphysema, those authors found a significant correlation between the quantitative assessment (with a computer algorithm) and the visual assessment. They also found that there was a considerable interobserver agreement in the visual assessment. Therefore, visual assessment of the distribution of pulmonary emphysema could be a reliable method, with one major advantage, which is the fact that everyone can use it, especially when CT analysis software is not available.

Our results suggest that, among COPD patients with emphysema, there is greater COPD severity, defined as a higher degree of airflow obstruction and lower alveolar-capillary diffusing capacity, in those with predominantly lower-lung emphysema, whereas functional status is better in those with predominantly upper-lung emphysema. These results can be explained, in part, by the smaller area of the lung affected when emphysema is predominantly in the upper lobes.

Regarding the COPD patients with homogeneous emphysema (type 3), our data indicate a tendency toward higher hyperinflation, with a higher RV/TLC ratio (Figure 2), although the difference did not reach statistical significance. That is probably associated with the broader, more uniform distribution of parenchymal destruction, together with the fact that had a median pack-year smoking history was higher among the patients with type 3 emphysema distribution (60 pack-years vs. 40.5 pack-years for those with type 1 emphysema distribution; \(p = 0.012\)).

After stratifying the study population according to the cut-off values for the assignment of at least 1 point on the **Body mass index, airflow Obstruction, Dyspnea, and Exercise capacity (BODE)** index,\(^\text{(21)}\) which assesses the risk of death for COPD patients, we observed that type 4 emphysema distribution (slightly predominant in the lower lung) significantly increases the risk of severe airflow obstruction (FEV\(_1\) < 65% of predicted) and reduced the 6MWD to < 350 m (Table 2). Hypoxemia was identified as another risk factor for mortality in COPD.\(^\text{(21,29)}\) Our results show that patients with emphysema that is slightly predominant in the lower lung are more likely to be hypoxemic.

Our findings are consistent with those of previous studies showing a strong association between lower-zone emphysema and airflow limitation.\(^\text{(18,19,30)}\) In another study, however, upper-zone predominance of emphysema was associated with a worse total St George’s Respiratory Questionnaire score, although it was not significantly associate with FEV\(_1\) (% of predicted).\(^\text{(31)}\) Reports are also inconsistent concerning the relationship between diffusing capacity and regional differences in emphysema distribution. Gurney et al.\(^\text{(32)}\) observed that DLCO is more strongly affected by lower-lung emphysema than by upper-lung emphysema, whereas Parr et al.\(^\text{(33)}\) found DLCO to be relatively preserved in patients with lower-lung emphysema. Those differences might be attributable to the different methods applied for assessing the regional distribution of emphysema.

The present study has a number of limitations. First, the female gender is not well represented in this study group. However, that is representative of the gender distribution of emphysema patients treated at our outpatient clinic. Second, our sample did not include any subjects with clearly lower lung-predominant emphysema (type 5). Because most of the patients were smokers, that type of emphysema distribution (sparing the upper lung) would be expected to be rather rare. We can presume that the clinical-radiological correlations for type 5 emphysema would be similar to those found for type 4. In order to extrapolate our results, a larger study sample, with similar gender proportions and including all types of emphysema distribution, will be needed. Finally, some interobserver

**Table 2. Distribution of pulmonary emphysema according to functional status and the respective functional severity.**

| Variable                  | Emphysema distribution | Frequency\( ^a \) n (%) | Adjusted OR\( ^b \) (95% CI) | p     |
|---------------------------|------------------------|-------------------------|-----------------------------|-------|
| FEV\(_1\) < 65% of predicted\(^d\) | Type 1                 | 14 (45.2%)              | 1 (reference)               | 0.045*|
|                           | Type 2                 | 13 (61.9%)              | 1.69 (0.32-8.92)            | 0.537 |
|                           | Type 3                 | 12 (85.7%)              | 5.79 (1.06-31.64)           | 0.043*|
|                           | Type 4                 | 15 (78.9%)              | 6.92 (1.43-33.45)           | 0.016*|
| 6MWD < 350 m\(^c\)        | Type 1                 | 1 (3.6%)                | 1 (reference)               | 0.064 |
|                           | Type 2                 | 2 (9.5%)                | 1.63 (0.31-8.70)            | 0.567 |
|                           | Type 3                 | 3 (21.4%)               | 5.58 (1.01-30.84)           | 0.049*|
|                           | Type 4                 | 9 (52.9%)               | 6.36 (1.26-32.18)           | 0.025*|
| Hypoxemia\(^d\)          | Type 1                 | 3 (13.0%)               | 1 (reference)               | 0.006*|
|                           | Type 2                 | 4 (20.0%)               | 2.85 (0.24-33.89)           | 0.408 |
|                           | Type 3                 | 6 (50.0%)               | 7.60 (0.67-86.19)           | 0.102 |
|                           | Type 4                 | 9 (52.9%)               | 32.66 (3.26-326.84)         | 0.003*|

6MWD: six-minute walk distance; type 1: obvious upper-lung predominance; type 2: slight upper-lung predominance; type 3: equal upper- and lower-lung extent; type 4: slight lower-lung predominance. *Corresponds only to patients with the lowest functional status, as defined in the first column. \(^a\)Adjusted for age and body mass index. \(^b\)Cut-off value for the assignment of at least 1 point on the Body mass index, airflow Obstruction, Dyspnea, and Exercise capacity (BODE) index. \(^c\)Defined as a PaO\(_2\) < 60 mmHg with an FiO\(_2\) of 0.21. \(^d\)p < 0.05.
variability is predictable, as previously noticed. Such disagreement can be seen primarily for patients with the least severe emphysema and with only partial upper or lower lung predominance. In fact, most discordant cases were related to classification differences between contiguous types.

In the past, direct visual observation and subjective visual grading were considered to have similar precision as the computer-assisted methods of emphysema quantification on CT scans. Although we have not provided a direct measure of emphysema severity, the purpose of this study was to present a qualitative (rather than quantitative), simple, affordable alternative method that could be widely used by clinicians to classify the heterogeneity of pulmonary emphysema.

In summary, in this group of COPD patients with pulmonary emphysema, lower lung-dominant distribution, as assessed by a subjective score, was found to have a significant impact on physiologic parameters, including pulmonary function test results and exercise capacity, although not on the clinical presentation of the disease, as assessed by the mRQC dyspnea scale score and the number of exacerbations in the last year. Further studies are warranted in order to confirm the importance of our findings.

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