Original paper

Lung density analysis using quantitative computed tomography in children with pectus excavatum

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Abstract

Purpose: To evaluate the mean lung density in children with pectus excavatum (PE) and to assess the correlation between the cardiac rotation angle, Haller index, pulmonary function test, and lung density.

Material and methods: This retrospective study included 33 children with PE and 31 healthy controls. The densities of lung parenchyma were evaluated by quantitative computed tomography (CT). Three lung levels were determined: T4 vertebra level, T10 vertebra level, and the level of the measurement of the cardiac rotation angle. The cardiac rotation angle and the Haller index were calculated. All measurements were done by 2 radiologists, independently. Student’s t-test or the Mann-Whitney U test, intraclass correlation coefficients, Pearson or Spearman’s rank correlation coefficient, and Kruskal-Wallis test were used for statistical analysis. A p-value less than 0.05 was considered as statistically significant.

Results: All the lung levels in the PE group had lower mean densities than healthy children, with statistical significance in the right lung at the T10 vertebra level (−818.60 ± 33.49 HU, −798.45 ± 40.24 HU; p = 0.028). There was a correlation between the cardiac rotation angle and the Haller index (r = 0.593; p < 0.001). There were no correlations between mean lung density and cardiac rotation angle, Haller index, and pulmonary function tests.

Conclusions: The lower mean lung densities were found in PE, especially in the right lower lobe. The parenchymal aeration should be considered independently from the severity of PE.

Key words: Haller index, lung density, pectus excavatum, paediatric, quantitative computed tomography.

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Authors’ contribution:

A Study design ∙ B Data collection ∙ C Statistical analysis ∙ D Data interpretation ∙ E Manuscript preparation ∙ F Literature search ∙ G Funds collection

Pectus excavatum (PE) is the most common congenital chest wall deformity with an incidence of 1/300–400 live births [1]. Displacement of the sternum and costosternal joints to posterior is the main occasion in PE. The condition often results in cardiopulmonary symptoms and psychological problems [2–5]. The most commonly associated anomalies are scoliosis, Marfan syndrome, congenital heart disease (most commonly mitral valve prolapse), rickets, Poland’s syndrome, and osteogenesis imperfecta [6].

Chest computed tomography (CT) is a key component to assess the mediastinal shift, cardiac compression, and the lung parenchymal abnormalities such as atelectasis or tracheobronchial compression in patients with PE. Indeed, subjective evaluations of the abnormal inflation that may be related to mediastinal shift and sternal depression on thorax CT scans have been found in PE patients [5,7]. However, the visual analysis may cause substantial variability in the subjective assessment among the readers. Quantitative CT analysis can reveal the exact
pixel densities of the entire lung or a specific area. Moreover, quantitative CT detects and calculates the percentage of abnormal parenchymal densities that would be extremely difficult to distinguish by visual analysis [8].

There have been several articles assessing lung parenchymal density in many acquired pulmonary diseases such as chronic obstructive pulmonary disease and interstitial lung disease, by using a sensitive density measurement method: quantitative CT [9-12]. Nonetheless, there is a lack of studies conducted by quantitative CT in congenital thoracic wall deformities even though the evaluation of parenchymal abnormalities in PE has recently become an area of interest [5].

We hypothesized that parenchymal hyperinflation may occur in patients with PE, especially in lower lung lobes. Thus, the purpose of this study is to evaluate the mean lung parenchymal density in children with PE comparing with healthy controls. The secondary goal is to assess the correlation between the cardiac rotation angle, Haller index, pulmonary function test, and parenchymal density in PE.

**Material and methods**

**Study population**

After receiving approval from Dokuz Eylul University School of Medicine institutional review board (2019/27-45), chest CT images of children with any thoracic wall deformity, who were admitted to the paediatric surgery department between January 2008 and March 2019, were retrospectively evaluated. The inclusion criteria were children between 0 and 18 years old with PE, and those with thorax CT imaging before PE repair surgery (n = 107). Exclusion criteria were children who had contrast-enhanced CT images (n = 25), alveolar or interstitial lung disease (n = 3), chemotherapy or radiotherapy history (n = 1), and images with motion artifacts (n = 31). Due to the extremely sensitive nature of quantitative CT analysis, children who had minimal artifacts in their chest CT were excluded to obtain accurate results. Normal mean values of the Haller index are described as less than 2.5 [13]. Patients with Haller index less than 2.5 were not accepted in the PE group (n = 14). After the inclusion and exclusion criteria, 33 patients were enrolled in this study.

As a control group, we scanned the chest CTs of paediatric patients who had thorax CT scanning between January 2008 and March 2019. Among these patients, only children with no abnormal findings on the CT scans were selected for the control group. The same exclusion criteria were used for the control group. Children with a Haller index higher than 2.5 who were detected incidentally were also excluded. Finally, 31 children were included in the control group. The parents of the children provided informed consent for imaging.

**Computed tomography protocols**

At our institution, CT scanning is routinely performed in full expiration for children. Verbal instructions and deep breathing exercises are performed before CT scanning. General anaesthesia or sedation is not performed while CT scanning.

CT images were acquired using a multi-detector CT scanner (Toshiba Aquilion 64; Toshiba Medical Systems, Otawara, Japan). Acquisitions were performed with no contrast injection and with the following parameters: 100 kV tube voltage, 50 mAs tube current, 3 mm slice thickness, and 0.6 mm collimation. CT dose index ranged from 0.67 to 7.5 mGy. Images were displayed with parenchymal 1600 Hounsfield units (HU) window width and −500 HU window level.

**Image interpretation**

**Quantitative lung density analysis**

Quantitative lung density analysis was performed using a semi-automated technique with the software package (Myrian 2.4.2., Intrasense, Montpellier, France). Three lung levels were determined for quantitative analysis of lung parenchymal density to observe the following: T4 vertebra spinous process level for upper lobes, T10 vertebra spinous process level for lower lobes, and the level at which the cardiac rotation angle was measured. The last level was determined to assess the effect of cardiac rotation on the lung parenchyma. All measurements were done for both lungs. Hereafter, we used “T4R” and “T4L” for the right lung and left lung at the T4 vertebra level, “T10R” and “T10L” for the right lung and left lung at the T10 vertebra level, and “CRAR” and “CRL” for the right lung and left lung at the cardiac rotation angle level. The T9 level was used when the lung parenchyma entering the slice at the T10 level was less than 75% of the total slice. The lung parenchyma was manually drawn on each slice to be the closest distance to the pleura (Figure 1). While drawing the medial border at the CRAR and CRAL levels, motion-induced densities were not included in the region of interest (ROI). Automated measurement of lung parenchymal density was obtained, and the parenchymal density distribution between −650 and −1050 was recorded as percentages.

**Cardiac rotation angle**

The cardiac rotation angle was accepted as the angle between the cardiac apex and the line between the anterior vertebral corpus and the spinous process (Figure 2).

**Haller index**

The Haller index was calculated on each participant utilizing the formula proposed by Haller et al. [13]: maximal trans-
verse diameter of the chest divided by the anterior-posterior distance on the axial slice, which demonstrates the smallest distance between the anterior surface of the vertebral body and the posterior surface of the sternum (Figure 2). The PE was severe if the Haller index was greater than 3.2 [14,15].

A radiologist with 8 years of experience and 3 years of experience in paediatric radiology, and a radiology resident with 3 years of experience performed the measurements individually. All measurements were performed twice by each radiologist.

Pulmonary function test

Spirometry (ZAN 510, Oberthulba, Germany) was performed according to the American Thoracic Society guidelines [16]. Spirometry provided the forced vital capacity (FVC), forced expiratory volume in 1 s (FEV1), and the FEV1/FVC ratio. The FVC and FEV1 were reported as the percentage predicted according to age, gender, and height. Normal spirometry findings were defined as FVC, FEV1, and FEV1/FVC ratio > 80% of the predicted value according to age, gender, and height. Obstructive lung function was defined as FEV1 < 80% of predicted value and FEV1/FVC < 80%. Restrictive lung function was described as a reduced FVC < 80% of predicted value, with a preserved FEV1/FVC ratio > 80%. The mean time between pulmonary function test administration and chest CT scanning was 7 days ± 11 (range, 0-50 days).

Statistical analysis

Statistical analyses were made using IBM SPSS ver. 22.0 (IBM Corp., Armonk, NY, USA). Categorical variables were summarized with frequency counts and percentages, and the continuous features were summarized with means and standard deviations. The distribution of the variables was performed using Kolmogorov-Smirnov and Shapiro-Wilk tests. Continuous variables were compared by using the Student's t-test or the Mann-Whitney U test. Interobserver agreements of the cardiac rotation angle, Haller index, and mean lung densities in each level were tested for intraclass correlation coefficients. The correlation between mean lung densities of each level and Haller index, cardiac rotation angle, and pulmonary function tests were measured with the Pearson or Spearman's rank correlation coefficient in the PE group. Correlation was considered slight when the correlation coefficient was < 0.2, fair when 0.2-0.4, moderate when 0.4-0.6, substantial when 0.6-0.8, and excellent when > 0.8. The Kruskal-Wallis test was used to compare the mean values of the patients according to the lung function tests. A p-value less than 0.05 was considered as statistically significant.

Results

In this study, 33 patients (girls/boys, 9/24; mean age, 14.42 years ± 2.25; age range, 10 to 18 years) and 31 healthy controls (girl/boy, 6/25; mean age, 14.61 years ± 2.09; age range, 10 to 18 years) were included. There were no significant differences in age and sex between the groups (p = 0.73; p = 1.0). The number of patients with Haller index < 3.2 was 13 (39.3%), while > 3.2 was 20 (60.6%). There were no patients who had a syndrome associated with PE. One patient had mild scoliosis. Demographic features and patients’ characteristics are shown in Table 1.
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Echocardiography revealed mitral valve prolapse in one patient with a Haller index of 3.05.

Excellent interobserver agreements were noted for quantitative analysis of the lungs (mean lung density values in each level, measurements of the Haller index and the cardiac rotation angle). The mean values of the 2 observers for each parameter were used in the final analyses because of the excellent agreement. The lowest mean lung density was found at the T4L level in the control group (mean, −801.23 ± 40.67 HU), T10R level in the PE (mean, −818.60 ± 33.49 HU). In comparison of mean lung densities of each level between two groups, significantly lower values were found at the T10R lung level in the PE group (−818.60 ± 33.49 HU, −798.45 ± 40.24 HU; \( p = 0.028 \)). The remaining levels did not show any significant differences regarding mean lung density, yet all the levels in PE group had lower mean lung densities (Table 2).

The cardiac rotation angle was measured as 46.71 ± 8.49° and 35.82 ± 5.38° in the PE group and control group, respectively (\( p < 0.001 \)). There was a significant correlation between the cardiac rotation angle and Haller index in the PE group (\( r = 0.593; \ p < 0.001 \)).

There were no correlations between any levels of the mean lung density and cardiac rotation angle and Haller index (Table 3). The mean lung density of the T10R lung level for children with Haller index < 3.2 was 815.84 ± 38.92 HU and for Haller index > 3.2 it was 820.40 ± 30.38 HU. The difference was not significant (\( p = 0.854 \)).

No significant differences were found between the sex groups and cardiac rotation angle and Haller index measurements in the PE group (\( p = 0.378; \ p = 0.725 \), respectively).

Pulmonary function tests were available in all PE patients. The spirometry results were normal in 21 of 33 patients (63.6%), obstructive pattern in 4 of 33 patients (12.1%), and restrictive pattern in 8 of 33 patients (24.2%). The mean values of the Haller index and the mean lung densities of each level did not show any significant difference between normal, obstructive, and restrictive pattern in pulmonary function tests (Table 4). No significant correlations were found between mean lung densities of each level and pulmonary function tests (Table 3).

**Discussion**

Lung parenchyma evaluation in patients with PE is important because PE causes not only cosmetic but also cardiopulmonary problems [2-5]. This is the first study evaluating the lung density of children with PE by a sensitive and objective method: quantitative CT. We demonstrated that all the lung levels had lower mean densities in the PE group compared to healthy children, with statistical significance in the right lung at the T10 vertebra level without any correlation with the Haller index, cardiac rotation angle, and pulmonary function tests. According to our results, we can speculate that the lower mean parenchymal densities may occur in children with PE regardless of the severity. Although most of the patients had normal pulmonary function test results, the lower mean lung densities on quantitative CT might indicate the initial phase of chronic lung changes.

Automated quantitative CT analysis provides more accurate results in the measurements of density, hetero-

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**Table 1. Patients’ characteristics, imaging, and clinical features**

<table>
<thead>
<tr>
<th>Factor</th>
<th>Pectus excavatum</th>
<th>Control</th>
<th>( p )-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age*</td>
<td>14.42 ± 2.25</td>
<td>14.61 ± 2.09</td>
<td>0.73</td>
</tr>
<tr>
<td>Sex (female/male)</td>
<td>9/24</td>
<td>6/25</td>
<td>1</td>
</tr>
<tr>
<td>Cardiac rotation angle*</td>
<td>46.71 ± 8.49</td>
<td>35.82 ± 5.38</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Haller index*</td>
<td>3.53 ± 0.66</td>
<td>2.05 ± 0.14</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Female*</td>
<td>3.39 ± 0.98</td>
<td>1.98 ± 0.12</td>
<td></td>
</tr>
<tr>
<td>Male*</td>
<td>3.56 ± 0.56</td>
<td>2.05 ± 0.13</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Pulmonary function tests</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>FEV1*</td>
<td>89.83 ± 10.32</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>FVC*</td>
<td>83 ± 10.90</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>FEV1/FVC*</td>
<td>105.26 ± 6.66</td>
<td>–</td>
<td></td>
</tr>
</tbody>
</table>

*Mean ± standard deviations. FVC – forced vital capacity, FEV1 – forced expiratory volume in 1 s

**Table 2. The comparison of the mean lung densities between the pectus excavatum and control groups**

<table>
<thead>
<tr>
<th>Mean lung density</th>
<th>Pectus excavatum</th>
<th>Control</th>
<th>( p )-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>T4R mean lung density</td>
<td>−814.75 ± 33.61</td>
<td>−793.19 ± 42.67</td>
<td>0.050</td>
</tr>
<tr>
<td>T4L mean lung density</td>
<td>−817.48 ± 39.66</td>
<td>−801.23 ± 40.67</td>
<td>0.094</td>
</tr>
<tr>
<td>T10R mean lung density</td>
<td>−818.60 ± 33.49</td>
<td>−798.45 ± 40.24</td>
<td>0.028</td>
</tr>
<tr>
<td>T10L mean lung density</td>
<td>−810.42 ± 39.25</td>
<td>−796.79 ± 41.89</td>
<td>0.202</td>
</tr>
<tr>
<td>CRAR mean lung density</td>
<td>−813.03 ± 40.02</td>
<td>−797.06 ± 36.33</td>
<td>0.087</td>
</tr>
<tr>
<td>CRAL mean lung density</td>
<td>−810.77 ± 41.72</td>
<td>−787.77 ± 39.44</td>
<td>0.060</td>
</tr>
</tbody>
</table>

T4R – the right lung at the T4 vertebra spinous process level, T4L – the left lung at the T4 vertebra spinous process level, T10R – the right lung at the T10 vertebra spinous process level, T10L – the left lung at the T10 vertebra spinous process level, CRAR – the right lung at the level of the measurement of cardiac rotation angle, CRAL – the left lung at the level of the measurement of cardiac rotation angle.
geneity, tissue, vascular, and airway by eliminating the intra- and interobserver variability [8]. There have been many studies regarding utilities of automated densitometric analysis by quantitative CT in adults [9-12]. However, only a few studies have been published including the use of an automated quantitative densitometric CT analysis in the paediatric age group [17-19]. Our study is the first article to calculate the lung parenchyma density by automated quantitative CT in children with PE.

Chest CT allows determination of several thoracic indexes to assess the severity of the chest wall deformity and to guide surgical strategies [19]. Although the degree of chest wall deformity is estimated objectively by the Haller index and correction index, the Haller index is commonly used to establish surgical candidacy [19]. Normal mean values of the Haller index are described as less than 2.5 [13]. The cut-off value of greater than 3.25 is defined for severe PE, and it is used as a surgical indicator according to the guidelines [14,15]. It is known that the phase of respiration at the time of imaging affects the Haller index [20,21]. We used inspiratory phase images in children because the cooperation for breath-holding was easier in children.

The mediastinal shift, cardiac compression, and parenchymal abnormalities can also be determined by chest CT. There have been several articles that have assessed sternal depression, cardiac rotation [7], Haller index [22], and parenchymal abnormalities [5,23] by using CT in patients with PE. In our study, the cardiac rotation angle increment was correlated with the Haller index in children with PE. The sternal depression may cause a right ventricular compression, and accordingly cardiovascular impairment may occur [24]. Chu et al. [7] reported that there was a positive correlation between the sternal depression index and the degree of cardiac rotation.

Figure 3. Histogram plots of the lung levels. T4R – the right lung at the T4 vertebra spinous process level; T4L – the left lung at the T4 vertebra spinous process level; T10R – the right lung at the T10 vertebra spinous process level; T10L – the left lung at the T10 vertebra spinous process level; CRAR – the right lung at the level of the measurement of cardiac rotation angle; CRAL – the left lung at the level of the measurement of cardiac rotation angle.
Singh et al. [5] proposed that both restriction of thorax anterior-posterior diameter and displacement of the heart to the left were the reasons for the compression of the bronchus and subsequently air being trapped in the lower lobe anterior basal segment. Our results did not support this theory. We suggest that lower mean lung densities might be associated with increased aeration, which may occur as a compensatory response to relatively pressed lungs due to the lower thoracic volume from birth. Also, the increased aeration in both sides of the upper and lower lobes may be explained by the fact that the disease is congenital.

Hyperinflation of the lower lobe anterior basal segment was reported in patients with PE in a study conducted by Singh et al. [5]. They also noted that this finding showed an association with the PE severity. They evaluated the presence or absence of left lower lobe hyperinflation by visual inspection and drawing manually a 1 cm² diameter ROI in the left and right anterior basal segments. It is known that the ROI provides information about the local density only in the selected area. As expected, focal hyperinflation could be observed by a small ROI that was placed on the visually selected focal parenchymal hypodense area. In our study, lower values of the mean parenchymal densities were noticed at all the levels, not at a focal area, in the PE group, with a statistical significance in the right lower lobe. Although our study contained a relatively small number of cases, we thought that we obtained more accurate results due to the sensitive nature of our measurement technique. The reason for not finding statistically significant lower results at all the levels in patients with PE may be the small study population. Additionally, the non-correlation between the mean lung densities and the Haller index in this study showed that the parenchymal hyperinflation develops independently of the severity of PE.

There are inconsistent results in the literature regarding whether the thoracic wall deformity in the PE causes an effect on pulmonary function [3–5,25–28]. Despite these controversies, pulmonary function tests are used as a surgical indicator and are performed for surgical candidates [15]. Most of the patients had normal spirometry results in our study (65.2%). Additionally, no significant relationships were found between the pulmonary function tests and mean lung densities in each level of the lung, as well as the Haller index severity and cardiac rotation angle. These
results may suggest that pulmonary function tests should be performed regardless of the severity of the Haller index and cardiac rotation in patients with PE. Our results were compatible with a study of Singh et al. [5]. Restrictive lung disease was the second most common spirometry finding in our study (21.7%). In a study by Lawson et al. [3] with a large cohort, it was reported that the restrictive pattern was more prominent than the obstructive pattern in PE patients. Additionally, the lung restriction was significantly related to the severity of the chest wall deformity. Non-correlated results between the pulmonary function tests and the Haller index in our study may be associated with the small number of patients in the group.

Our study has some limitations. First, we examined a relatively small sample size. Although many patients' images had enough quality for preoperative evaluation, all the patients who had minor artifacts were excluded for reliable and accurate densitometric analysis. Second, the lung density and the Haller index are related to the phase of breathing on chest CT scans. Inspiratory images alone are unlikely to be the best option to check the effect of PE on chest mechanics. However, as in our hospital and in many centres, chest CT scans are done with a single phase routinely in children to minimize the effects of the radiation.

**Conclusions**

This study demonstrated lower mean lung densities in children with PE compared to healthy children, especially in the right lower lobe. However, the lower mean lung densities were not associated with either the Haller index or cardiac rotation. Pulmonary function tests were also not correlated with mean lung densities. We recommend considering the parenchymal aeration independently from the severity of PE. Finally, further prospective research regarding the potential clinical significance and long-term assessment is needed because there may be cardiopulmonary consequences of the decreased mean lung density.

**Conflict of interest**

The authors report no conflict of interest.

**References**


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