

RESEARCH

Open Access

Low-dose CT measurements of airway dimensions and emphysema associated with airflow limitation in heavy smokers: a cross sectional study

Akkelies E Dijkstra^{1*}, Dirkje S Postma¹, Nick ten Hacken¹, Judith M Vonk^{1,2}, Matthijs Oudkerk³, Peter MA van Ooijen³, Pieter Zanen⁴, Firdaus A Mohamed Hoesein⁴, Bram van Ginneken⁵, Michael Schmidt⁶ and Harry JM Groen¹

Abstract

Background: Increased airway wall thickness (AWT) and parenchymal lung destruction both contribute to airflow limitation. Advances in computed tomography (CT) post-processing imaging allow to quantify these features. The aim of this Dutch population study is to assess the relationships between AWT, lung function, emphysema and respiratory symptoms.

Methods: AWT and emphysema were assessed by low-dose CT in 500 male heavy smokers, randomly selected from a lung cancer screening population. AWT was measured in each lung lobe in cross-sectionally reformatted images with an automated imaging program at locations with an internal diameter of 3.5 mm, and validated in smaller cohorts of patients. The 15th percentile method (Perc15) was used to assess the severity of emphysema. Information about respiratory symptoms and smoking behavior was collected by questionnaires and lung function by spirometry.

Results: Median AWT in airways with an internal diameter of 3.5 mm (AWT_{3.5}) was 0.57 (0.44 - 0.74) mm. Median AWT in subjects without symptoms was 0.52 (0.41-0.66) and in those with dyspnea and/or wheezing 0.65 (0.52-0.81) mm ($p < 0.001$). In the multivariate analysis only AWT_{3.5} and emphysema independently explained 31.1% and 9.5% of the variance in FEV₁% predicted, respectively, after adjustment for smoking behavior.

Conclusions: Post processing standardization of airway wall measurements provides a reliable and useful method to assess airway wall thickness. Increased airway wall thickness contributes more to airflow limitation than emphysema in a smoking male population even after adjustment for smoking behavior.

Keywords: Airway dimensions, Low-dose CT, Respiratory symptoms, Smoking, Airflow limitation, Emphysema

Introduction

The quantification of airway dimensions by CT has become feasible with the development of multi detector computed tomography (CT) and new software tools for image analysis [1,2]. Assessment of airway dimensions by CT has been studied particularly in relation to

asthma, smoking and chronic obstructive pulmonary disease (COPD) [3-8], diseases generally associated with chronic or intermittent airflow limitation. So far, airway wall thickness (AWT) measurements have been performed by selecting well quantifiable airways [9-12] or by standardizing dimensions to airways with a 10 mm internal lumen perimeter (pi10, equivalent to about 3.2 mm internal airway lumen diameter) derived from a small number of airways [5,6,13-15].

More recently, low-dose multi slice CT has become available, a technique with a good quantifying performance

* Correspondence: a.e.dijkstra@umcg.nl

¹University of Groningen, Department of Pulmonary Diseases, University Medical Center Groningen, GRIAC research institute, Groningen, the Netherlands

Full list of author information is available at the end of the article

that is preferred for monitoring of pulmonary and airway abnormalities as compared to the high radiation exposures with high-resolution computed tomography (high-dose CT). The cumulative radiation dose exposure with low-dose CT remains very low, even when individuals are exposed multiple times [16]. Airway dimensions measured with low-dose CT have only been reported in few studies using the same diversity in analytic approaches as applied with high-dose CT measurements [7,17-19].

Airway dimensions and extent of emphysema are known to be associated with airflow limitation [6,9-14,19,20], although the influence of smoking behavior or signs of airway disease such as cough, dyspnea, wheezing or mucus overproduction on airflow limitation is not clear [5,7,21,22].

The aims of this study are to quantify airway dimensions of the lung in multiple airway sections of each lobe in a novel manner and the extent of emphysema by using low-dose CT. These measurements were associated with the influence of airflow limitation, respiratory symptoms and corrected for smoking behavior.

Methods

Population

We randomly selected 500 current and former smokers participating in the Groningen cohort of a male population-based multi-centre lung cancer screening study (NELSON). The Dutch ministry of health and the Medical Ethics Committee of the hospital approved the study protocol. Informed consent was obtained from all participants. Detailed inclusion criteria and characteristics have been described elsewhere [23]. In short, subjects with a smoking history of at least 20 pack-years were included. Information on the presence or absence of respiratory symptoms and smoking (pack-years and former or current smoking) was obtained by questionnaires. The question used to record respiratory symptoms was "do you have experienced the following symptoms cough, sputum expectoration, wheezing or dyspnea for at least 3 months during the past year, even when you did not have a cold?"

Lung function

All participants performed standardized spirometry according to the European Respiratory Society guidelines [24], including forced expiratory volume in 1 sec (FEV₁) and forced vital capacity (FVC) at the start of the study. In this population-based study we did not administer a bronchodilator.

CT scanning

Low-dose CT images of the chest were acquired at full inspiration after appropriate instruction on one CT scanner (Sensation 16, Siemens Medical Solutions, Forchheim, Germany) [23] according to the following

protocol: spiral acquisition at 120 kV, 20mAs, rotation time 0.5 s, pitch 1.5 and collimation 16×0.6 mm, field of view 300 to 350 mm, slice thickness 1 mm and slice increment 0.7 mm. The effective radiation dose was less than 0.8 mSv. Contrast medium was not used. The images were reconstructed to a pixel matrix of 512×512 using B30f kernel. Thus, the spatial resolution was 0.59 to 0.68 mm in *x-y* plane, and 0.7 mm in *z* plane. The CT system was calibrated routinely.

Quantification of AWT

AWT was measured in cross-sectionally reformatted images with an automated research software prototype MEVIS Airway Examiner v1.0 (release 2009, Fraunhofer MEVIS, Bremen, Germany) based on an algorithm by Weinheimer at locations with a fixed internal diameter of 3.5 mm in each lung lobe [25]. This software automatically extracts airway centerlines, re-samples images perpendicular to the airway direction at equally spaced positions along the centerline and detects inner and outer airway wall borders in these images. The outer wall border is detectable when no adjacent tissue with similar CT density is present and is taken into account when the wall is detected in at least 25% of the perimeter at a location. AWT and the fraction of perimeter where the outer wall border was identified (Assessed Perimeter Fraction, APF) are calculated for each location. Wall thickness quantification accounts for partial volume effects by integrating Hounsfield units across the wall. Accuracy and reproducibility of this algorithm was tested previously under clinical conditions using a similar protocol as used in our study [2]. Average wall thickness and cumulative APF of all detectable airway locations with a fixed lumen diameter is reported per lobe and for the whole lung. The borders of the lung lobes were automatically calculated by the software in a standard way. All low-dose CT scans were visually evaluated for appropriate segmentation.

Quantification of emphysema and lung volume

Quantification of emphysema was based on density differences and measured with a software tool called Image Explorer (Image Sciences Institute, Utrecht, the Netherlands) [16,26]. This software produces automatically the lung volume. The extent of emphysema was automatically performed at the 15th percentile (Perc15) of the Hounsfield density distribution. Perc15 is the threshold density value where 15% of all voxels has a lower density [27]. A lower Perc15, i.e. closer to -1000 HU, means that more emphysema is present. All scans were reconstructed with a soft reconstruction filter (Philips B, Siemens B30f). Airways were automatically excluded to assess density of lung parenchyma exclusively and HU densities of the entire scan were recalibrated using automatically measured average

densities in the trachea and shifting the HU values of the entire scan so that air density in the trachea became -1000 HU. Additionally, the percentage of low attenuation area, defined as the proportion of low-density voxels below -950 HU (%LAA-950HU) was used. %LAA-950HU was log-transformed because of skewed distribution.

Explorative studies

Prior to the research described above we have

- 1) established the optimal internal airway diameter, i.e. the internal airway diameter that allows the highest number of cumulatively Assessed Perimeter Fractions (APF) for the whole lung. Therefore we measured APF on 20 selected NELSON CT's in airways with a lumen diameter of 2.5, 3.0, 3.5, 4.0, 4.5 and 5.0 mm (± 0.25 mm) divided into 3 groups: no emphysema and normal lung function ($n = 8$, $p^{15} > -920$ and $FEV_1/FVC > 85\%$), moderate emphysema and normal lung function ($n = 4$, $-940 < p^{15} < -960$ and $FEV_1/FVC > 70\%$) and no emphysema and severe airflow limitation ($n = 8$, $p^{15} > -920$ and $FEV_1/FVC < 50\%$).
- 2) compared the mean $AWT_{3.5}$, using the same method as described above, at the established internal lumen size with high- and low-dose CT in 8 NELSON subjects from whom high- and low-dose CT were available. These CT data were obtained in spiral mode with 16×0.75 mm collimation and in full inspiration with the same scanner (Sensation-16 Siemens Medical Solutions, Forchheim, Germany). Axial images were reconstructed with 1.0 mm thickness at 0.7 mm increments. All scans were reconstructed with a soft reconstruction filter (Siemens B30f) at a 512×512 matrix.
- 3) determined the generation where airways with the established optimal internal lumen size are present. AWT measurements at 3.5 mm internal lumen size were performed in 57 randomly selected low-dose CTs of NELSON subjects. A multi-planar reconstruction (MPR) was made in each case of the apical upper lobe bronchus (B1) and the posterior lower lobe bronchus (B10). Subsequently the location was projected on the segmentation image. Three-dimensional image moving created the opportunity to observe airways from various directions and to check bifurcations and count airway generations according to the method of Boyden [28].

Statistical analysis

Data are reported as mean \pm standard deviation (SD) or median (25th - 75th percentile) values as appropriate. The mean AWT at 3.5 mm internal lumen size

($AWT_{3.5}$) of all five lobes per case was calculated taking into account the APF per lobe by the following formulae: $((AWT \text{ left upper lobe} \times APF \text{ left upper lobe}) + (AWT \text{ left lower lobe} \times APF \text{ left lower lobe}) + (AWT \text{ right upper lobe} \times APF \text{ right upper lobe}) + (AWT \text{ right middle lobe} \times APF \text{ right middle lobe}) + (AWT \text{ left upper lobe} \times APF \text{ left upper lobe})) / \text{sum of APF of all lung lobes}$. $AWT_{3.5}$ for the whole population was skewed distributed, therefore we report median AWT and range, and log-transformed AWT was used in the analyses.

Differences between various categories were explored using chi-square tests (dichotomous data), 2-tailed unpaired Student's t-tests for normally distributed continuous data and Mann-Whitney U-tests for not normally distributed continuous data. The difference in airway wall thickness between lung lobes was assessed with a Wilcoxon signed rank test. Univariate linear regression analyses was used to study the relationships between clinical variables and AWT, and those variables with $FEV_1\%$ predicted. Next, multivariate linear regression analyses were performed on those clinical variables showing significance in the univariate regression analyses. Outcomes of these analyses have been described with beta's and p-values. Bland-Altman plot was used to analyze the agreement between AWT by high- and low-dose CT [29]. All statistical analyses were performed using SPSS 18.0 for Windows; P-values below 0.05 were considered statistically significant.

Results

Population characteristics

After visual evaluation 8 out of the 500 randomly selected subjects were excluded because of (partial) missing of airway segmentation on CT. The mean age of the cohort was $59.4 (\pm 5.2)$ years, approximately 59% were current smokers and median pack-years was 34.0 (28.0 - 45.6). More than 51% of the population reported at least one respiratory symptom (Table 1).

Airway wall thickness

Explorative analyses

The highest numbers of cumulatively assessed perimeter fractions (APF) of airways were reached at an internal lumen perimeter of 3.5 ± 0.25 mm (Figure 1); therefore this diameter was selected for further analysis. Median $AWT_{3.5}$ on low-dose CT was comparable with median $AWT_{3.5}$ on high-dose CT, respectively 0.57 (0.48 - 0.74) mm and 0.55 (0.47 - 0.73) mm ($p = 0.89$, $n = 8$). This demonstrates that MEVIS software analysis of data from low-dose CT gives similar results as from high-dose CT (Figure 2). Airways with an internal diameter of 3.5 mm appeared mainly in the 5th generation (range, 3rd - 7th generation) in the upper lobe bronchus, and mainly in the 8th - 10th generation (range, 6th - 12th generation) in

Table 1 Clinical and demographic characteristics of heavy smokers from the general population cohort

N	492
Age, years	59.4 ± 5.2
Pack-years smoking	34.0 (28.0 - 45.6)
Current smoking,%	59.1
FEV ₁ , liter	3.45 ± 0.76
FEV ₁ ,%predicted	98.2 ± 19.7
FEV ₁ /FVC,%	70.0 ± 10.7
Emphysema; Perc15, HU	-920 (-930 to -907)
Emphysema;%LAA -950 HU	2.5 (1.3 - 4.3)
Emphysema;>5%LAA -950 HU,%	19.7
Lung volume on CT, liter	6.5 ± 1.4
Chronic Mucus Hypersecretion,%	29.7
Cough,%	32.7
Dyspnea,%	28.4
Wheezing,%	25.3
No respiratory symptoms,%	48.6

Mean ± standard deviation shown for continuous data and median (interquartile range) for non-parametric distribution.

Definition of abbreviations: HU = Hounsfield Units; Perc15 = the threshold density value where 15% of all voxels has a lower density; %LAA -950 HU = percentage low attenuation areas < -950 HU; >5%LAA -950 HU,% = percentage of the population having >5% low attenuation areas < -950 HU.

the lower lobe bronchus. This distribution was observed irrespective of smoking, presence of airflow limitation (defined as FEV₁/FVC < 0.70) or chronic mucus hypersecretion (CMH) (Figure 3).

Airway wall thickness in the population

In the whole population the median (25th - 75th percentile) AWT_{3.5} was 0.57 (0.44 - 0.74) mm. The APF in the whole lung varied from 142 to 295 (median 215). The results per lung lobe are presented in the online Additional file 1: Table S1).

Airway wall thickness and clinical characteristics

Significantly higher AWT_{3.5} values were observed in subjects with dyspnea and/or wheezing (n = 181, median AWT_{3.5} 0.66 mm) or with cough and/or CMH (n = 201, median AWT_{3.5} 0.63 mm) compared to subjects without dyspnea and/or wheezing (n = 309, median AWT_{3.5} 0.53 mm, p < 0.001) or without cough and/or CMH (n = 291, median AWT 0.53 mm, p-values < 0.001). Current smokers and former smokers had comparable median AWT_{3.5} values, i.e. 0.58 mm and 0.56 mm (p = 0.113) respectively (Table 2).

Univariate linear regression analysis showed inverse associations between log-AWT_{3.5} and FEV₁ (b = -0.233, p < 0.001), FEV₁/FVC (b = -0.015, p < 0.001), FEV₁% predicted (b = -0.010, p < 0.001) and lung volume (b = -0.055, p < 0.001) and positive associations between

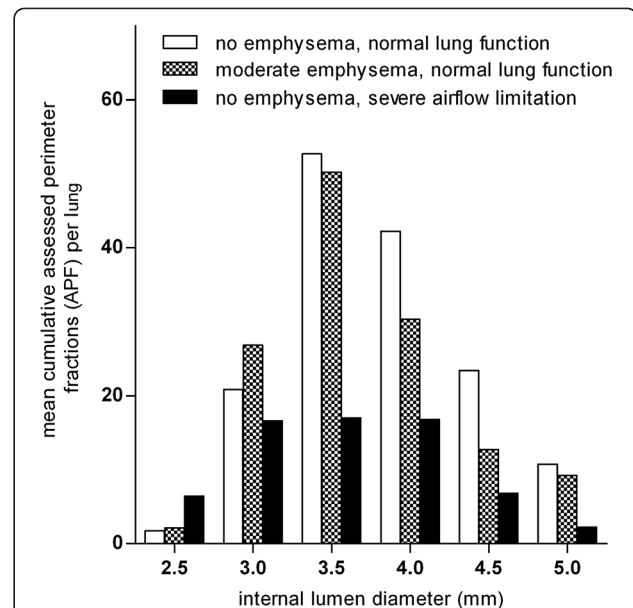


Figure 1 Determination of the optimal airway size. Mean cumulative assessed perimeter fractions in the total lung, for different groups of patients at different internal airway lumen diameters. APF was measured on low-dose CT of 20 selected NELSON subjects divided into 3 groups: subjects without emphysema and with normal lung function (n = 8, perc15 > -910 and FEV₁/FVC > 85 %), with moderate emphysema and normal lung function (n = 4, -940 < perc15 < -960 and FEV₁/FVC > 70 %) and without emphysema and having severe airflow limitation (n = 8, perc15 > -920 and FEV₁/FVC < 50 %), in airways with a lumen diameter of 2.5, 3.0, 3.5, 4.0, 4.5 and 5.0 mm (± 0.25 mm).

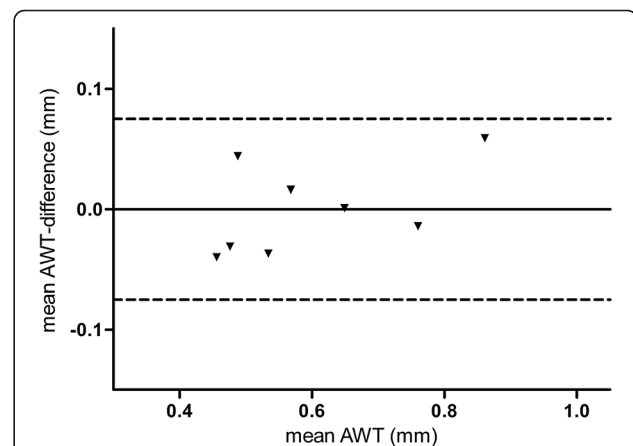


Figure 2 Comparison of airway wall thickness on high- and low-dose CT. AWT_{3.5} was measured on high and low-dose CT of 8 NELSON subjects. Bland & Altman plot shows agreement between mean AWT_{3.5} determined by high- and low-dose CT. Dashed lines depict the 95% confidence interval.

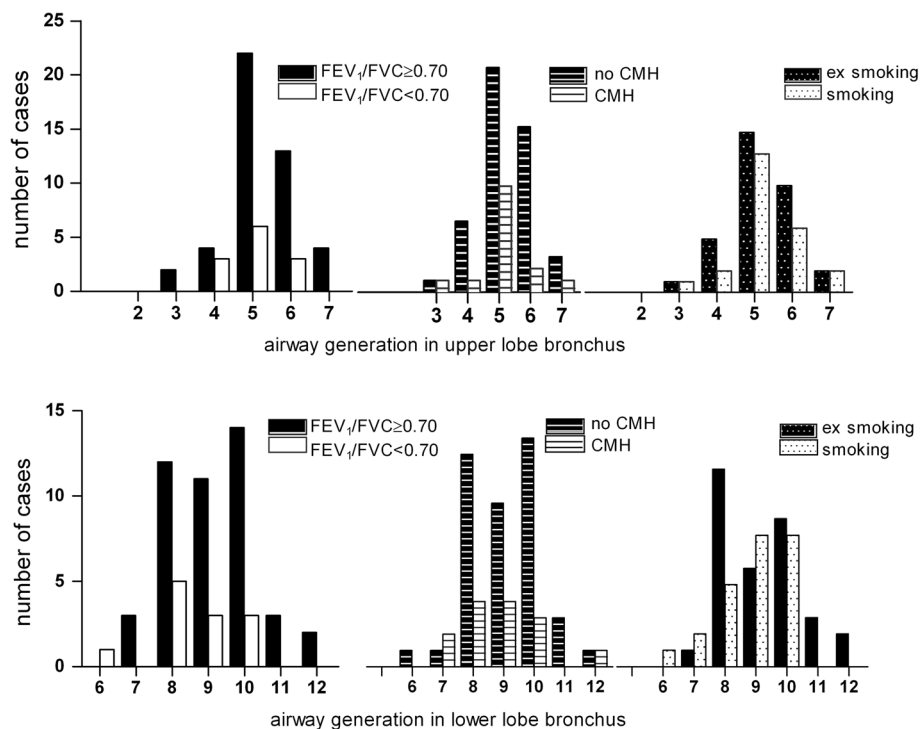


Figure 3 Distribution of 3.5 mm sized airways over airway generations. The distribution of 3.5 mm internal lumen sized airways over the 2nd till 7th airway generations in the apical upper lobe bronchus (B1) and over the 6th till 12th generation in the lower lobe bronchus (B10) in the right lung, for subjects with and without airway obstruction, for subjects with and without Chronic Mucus Hypersecretion (CMH) and for current and former smokers. The number of cases (e.g. no CMH and CMH) in each group was limited but the distribution was similar.

log-AWT_{3.5}, Perc15 and number of pack-years smoking, respectively ($b = 0.003$, $p < 0.001$) and ($b = 0.003$, $p = 0.003$) (Table 3).

Multivariate analysis on all clinical variables significantly associated with AWT in univariate analyses revealed that log-AWT_{3.5} was independently associated with lower FEV₁%predicted ($b = -0.010$, $p < 0.001$), higher Perc15 ($b = 0.005$, $p < 0.001$), and lung volume ($b = -0.037$, $p = 0.005$) respectively (Table 4).

Contribution of airway wall thickness and emphysema to airflow limitation

To study the contribution of AWT_{3.5}, emphysema (Perc15), and clinical variables to airflow limitation, FEV₁%predicted was taken as dependent variable. A significant negative association was found between FEV₁%predicted and log-AWT_{3.5} ($b = -31.21$, $p < 0.001$), pack-years ($b = -0.153$, $p = 0.004$), cough, CMH, dyspnea and wheezing ($b = -9.234$, -8.17 , -11.99 , -13.26 respectively, all p -values < 0.001), and a positive association between

Table 2 Airway wall thickness in subjects with and without respiratory symptoms and in current and former smokers

	n	AWT _{3.5} (mm)		n	AWT _{3.5} (mm)	p-value
CMH	146	0.62 (0.49-0.80)	no CMH	346	0.56 (0.44-0.71)	0.002*
Cough	161	0.64 (0.50-0.84)	no cough	331	0.54 (0.43-0.70)	<0.001*
Dyspnea	140	0.65 (0.51-0.80)	no dyspnea	352	0.54 (0.43-0.69)	<0.001*
Wheezing	125	0.66 (0.52-0.85)	no wheezing	367	0.53 (0.43-0.70)	<0.001*
CMH and/or cough	201	0.63 (0.49-0.79)	no CMH or cough	291	0.53 (0.43-0.70)	<0.001*
Dyspnea and/or wheezing	181	0.66 (0.52-0.83)	no dyspnea or wheezing	311	0.53 (0.42-0.67)	<0.001*
Cough, CMH, dyspnea and wheezing	49	0.69 (0.51-0.89)	no cough, CMH, dyspnea or wheezing	239	0.52 (0.41-0.66)	<0.001*
Current smoking	291	0.58 (0.45-0.75)	Former smoking	201	0.56 (0.43-0.71)	0.113

Data presented as median (interquartile range) values.

Definition of abbreviations: AWT_{3.5} = airway wall thickness at 3.5 mm internal lumen size; CMH = chronic mucus hypersecretion.

Table 3 Univariate associations between (A) log-transformed AWT_{3.5} and (B) FEV₁% predicted, and clinical characteristics

Dependent variable	A. Log-AWT _{3.5}		B. FEV ₁ % predicted	
	Beta	p-value	Beta	p-value
Log-AWT _{3.5}			-31.21	<0.001
FEV ₁ % predicted	-0.010	<0.001		
FEV ₁ , liter	-0.233	<0.001	23.56	<0.001
FEV ₁ /FVC,%	-0.015	<0.001	1.441	<0.001
FVC,% predicted	-0.009	<0.001	0.971	<0.001
Emphysema; Perc15	0.003	<0.001	0.204	<0.001
Emphysema; Log-%LAA -950 HU	-0.041	0.023	-5.872	<0.001
Lung volume on CT, Liter	-0.055	<0.001	-0.257	0.691
Pack-years	0.003	0.003	-0.153	0.004
Smoking (former/current)	0.059	0.065	-3.257	0.071
Age, years	-0.002	0.458	-0.289	0.092
Height, cm	-0.002	0.444	-0.121	0.425
Cough	0.156	<0.001	-9.234	<0.001
CMH	0.114	0.001	-8.171	<0.001
Dyspnea	0.158	<0.001	-11.99	<0.001
Wheezing	0.202	<0.001	-13.26	<0.001

Definition of abbreviations: Log-AWT_{3.5} = log transformed airway wall thickness at 3.5 mm diameter; CMH = chronic mucus hypersecretion; Log-%LAA -950 HU = log transformed percentage of low attenuation areas < -950 HU.

FEV₁%predicted and Perc15 (b = 0.204, p < 0.001) (Table 3).

Multivariate analysis on all clinical variables that were associated with FEV₁%predicted in univariate analyses showed that higher FEV₁%predicted was significantly associated with lower AWT_{3.5} values (b = -31.3, p < 0.001) and with higher Perc15 (b = 0.342, p < 0.001) (Table 4). AWT_{3.5} and Perc15 explained 31.1% and 9.5% of the variance of FEV₁%predicted, respectively. The

Table 4 Multivariate linear regression: dependent variable is (A) log transformed AWT_{3.5} and (B) FEV₁% predicted

Dependent variable	A. Log-AWT _{3.5}		B. FEV ₁ % predicted	
	Beta	p-value	Beta	p-value
FEV ₁ % predicted	-0.010	<0.001		
Log-AWT _{3.5}			-31.28	<0.001
Emphysema; Perc15	0.005	<0.001	0.342	<0.001
Lung volume	-0.037	0.005	0.209	0.773
Pack-years	0.002	0.031	-0.029	0.499
Smoking (former/current)	-0.011	0.700	-2.756	0.086
Cough	0.030	0.404	-0.408	0.834
CMH	-0.026	0.447	-2.098	0.262
Dyspnea	-0.007	0.828	-2.945	0.116
Wheezing	0.058	0.124	-4.004	0.051

Definition of abbreviations: Log-AWT_{3.5} = log transformed airway wall thickness at 3.5 mm diameter; CMH = chronic mucus hypersecretion.

results of the multivariate regression analysis with emphysema expressed as %LAA-950 HU as independent variable, are presented in the online Additional file 1: Table S2.

Discussion

Low-dose CT is an appealing approach to quantify simultaneously pulmonary and airway abnormalities. Our study shows that the use of low-dose CT combined with modern post processing software tools provides reliable information on airway wall thickness and the extent of emphysema in a heavy smoking male population. Although CT does not provide dynamic measurements, airway wall thickening and emphysema explained respectively 31.1% and 9.5% of the variance in FEV₁%predicted, the most commonly used variable of airflow limitation. Changes in AWT of more than 0.1 mm reflecting lumen surface changes over 8% measured at one air lumen level were observed between cases with and without respiratory symptoms.

Our study confirmed that increased AWT is associated with lower FEV₁%predicted. This lower FEV₁%predicted depends on the airway size in which the measurements of AWT are being performed [12,17] and on the characteristics of the study population [7,20]. Our population consisted of rather healthy elderly males from a randomly recruited Dutch heavy smoking population and still we were able to find significant associations between

thicker airway walls and more severe airflow limitation. In contrast with the study of Nakano we found a significant negative association between AWT and FEV₁/FVC illustrating the sensitivity of our method [9].

The significant but weak negative association between airway wall thickness and emphysema has also been reported in other studies [7,13,30] but was not found in the study by Nakano [9]. Loss of elastic recoil may contribute to collapse of the airways resulting in a more proximal localization of airways with 3.5 mm internal lumen diameter. As these more proximal airways have thicker airway walls this phenomenon contributes to the weak negative association. Another possible explanation for this negative association may be that there are subjects with predominantly airway wall thickening and others with predominantly emphysematous changes. Particularly subjects with relatively more airway wall thickening are responsible for the negative association and subjects with predominantly emphysematous changes do hardly contribute. Apparently, in our population of subjects with normal lung function and with mild airflow limitation, the bronchitic phenotype is already present in the very early stages of smoking-induced lung disease. Discrepancies between the study of Nakano [9] and our study may be due to the composition and size of the study populations, respectively predominantly emphysema versus predominantly healthy smokers with respiratory symptoms.

Importantly, we observed that the contribution of AWT_{3.5} to airflow limitation was larger than the emphysema component. Moreover, AWT_{3.5} and emphysema together only explained about 40% of the variance in FEV₁%predicted in this smoking male population. This unexpected low contribution of AWT_{3.5} and emphysema to FEV₁%predicted may be that the CT images were obtained at full inspiration, while FEV₁ reflects expiratory airflow limitation. One explanation for this observation is that airflow limitation is not only due to reduced airway diameter at one level but should be evaluated as an integral of all airways at all lumen diameters. This is difficult to achieve and therefore we took the smallest measurable lumen diameter that provides the largest contribution to airflow limitation. A more obvious physiological explanation may be the presence of the heterogeneity in airway ventilation interrupting the symmetry in parallel airways leading to large clusters of poorly ventilated lung units [31].

In the univariate analysis, increased AWT_{3.5} was associated with respiratory symptoms. However, AWT_{3.5} was not associated with the presence of any respiratory symptom in the multivariate analysis after adjustment for FEV₁%predicted, emphysema and smoking behavior. This finding corresponds with other studies [7] and is consistent with the idea that inflammation and airway

remodeling, associated with chronic bronchitis, is located in the more central airways [32]. The study of Martinez et al. showed a positive association between airway dimensions and questionnaires, the BODE index [33] and the St. George's respiratory questionnaire [34] including questions about BMI, respiratory symptoms, exercise capacity and lung function. Also Camiciottoli et al. found a positive association between BODE and airway wall thickness [35]. Our study also showed that including respiratory symptoms in the multivariate model with AWT_{3.5} and emphysema has no impact on airflow limitation.

Lung volume depends on height, weight and sex and as a consequence each person has different airway dimensions. Therefore, airway dimensions should be corrected for lung volume. Actually, volume-corrected AWT is the best parameter to use. In this study lung volume does not change the multivariate model because FEV₁%predicted is already corrected for lung volume by correcting for patient height.

It has been shown that the automated imaging program (MEVIS Airway Examiner) based on a method by Weinheimer et al. performed much better than the often used "full-width-at-half-maximum" method in a silicon tube phantom, regarding the blurring effect of CT [25,36]. Usually it is better to use sharper kernels for airway quantification. However, it was shown in the study by Schmidt et al. that the MEVIS airway examiner provides reproducible quantitative results across different reconstruction kernels (B30f and B50f) and repeated acquisitions [2]. Moreover, the "full-width-at-half-maximum" technique systematically overestimates AWT, particularly in small airways [36]. Because low-dose CT and the automated imaging program (MEVIS Airway Examiner) had not been used previously in clinical studies, we firstly optimized our post processing measurements in smaller cohorts of patients before applying it in the population study. We demonstrated that the highest number of AWT measurements could be performed on airways with an internal diameter of 3.5 mm, irrespective of the presence of airflow limitation or emphysema. In addition we demonstrated that differences in AWT_{3.5} are not explained by differences in airway generations. Finally, we demonstrated that low-dose CT imaging provided similar AWT results as high-dose CT imaging.

In a non-biased way we were able to evaluate 230 cumulatively assessed perimeter fractions (APF) per CT, ranging from 27-641 APF. In contrast to the commonly used pi10 method, in which a secondary derived variable from few, mostly 6 selected airways is used to estimate the airway wall thickness [5,6,13-15], we obtained many direct airway wall measurements. To our opinion direct measurements assessed over all lobes provide a better overall reflection of AWT than a limited number of secondary AWT measurements.

A limitation of this study is, inherent to general population-based studies, that only male smokers with mild emphysema and/or airflow limitation were included. Strengths of our study is the non-biased way of analyzing a high number of airway sections in all lobes that makes our approach more suitable for combined airway wall thickness and emphysema measurements on one low-dose CT scan. Such approach allows monitoring of intervention effects on both parameters. This is important when new treatment modalities will become available for clinical testing.

In the future further developments may involve measurements of thickness of airway walls along the full length of the bronchial tree at in- and expiration scans. Possibly more airflow variability will be explained.

In conclusion, post processing standardization of large numbers of airway wall measurements in all lung lobes is feasible, reliable and an useful method to assess airway wall thickness. We have demonstrated that increased airway wall thickness contributes more to airflow limitation than emphysema in a smoking male population even after adjustment for smoking behavior.

Additional file

Additional file 1: Table S1. Median $AWT_{3.5}$ and cumulatively assessed perimeter fractions (APF) per lung lobe. **Table S2.** Multivariate linear regression: dependent variable is (A) log transformed $AWT_{3.5}$ and (B) $FEV_1\%$ predicted. $LAA\% \geq 950$ HU was used to quantify the extent of emphysema in this analysis.

Abbreviations

APF: Cumulative assessed perimeter fractions; AWT: Airway wall thickness; $AWT_{3.5}$: Airway wall thickness at 3.5 mm internal lumen diameter; BMI: Body mass index; CMH: Chronic mucus hypersecretion; CT: Computed tomography; FEV_1 : Forced expiratory volume in 1 sec; FEV_1/FVC : Ratio of forced expiratory volume in 1 sec and forced vital capacity; $FEV_1\%$ predicted: (Ratio of present FEV_1 and expected FEV_1) \times 100 %; FVC: Forced vital capacity; GOLD: the Global Initiative for Chronic Obstructive Lung Disease; HRCT: High resolution computed tomography; HU: Hounsfield unit; %LAA: Percentage of low attenuation areas; NELSON: the Dutch-Belgian Randomized Lung Cancer Screening Trial; Perc15: the threshold density value where 15% of all voxels has a lower density; PFT: Pulmonary function test; SD: Standard deviation.

Competing interests

Dr Postma received funds for research from AstraZeneca, Chiesi, GSK. DSP has been consultant to AstraZeneca, Boehringer Ingelheim, Chiesi, GSK, Nycomed and TEVA. Travel to meetings has been funded by AstraZeneca, Chiesi, GSK and Nycomed. Dr Zanen received an EU FP7 grant. MSc Schmidt received a grant for development of algorithms for the airway examiner by MEVIS Medical Solutions. MSc Dijkstra, Dr ten Hacken, Dr Vonk, Dr Oudkerk, Dr van Ooijen, Dr Mohamed Hoesein, Dr van Ginneken and Dr Groen have reported that no potential conflicts of interest exist with any companies/organizations relevant to this article.

Authors' contributions

AD designed the study concept, collected data, performed statistical analysis, interpreted data and drafted, wrote and finalized the manuscript; HG and DP designed the study concept, collected data, interpreted data and drafted and finalized the manuscript; NH and FMH interpreted data and drafted and finalized the manuscript; JV performed statistical analysis, interpreted data

and drafted and finalized the manuscript; MS designed and supervised the use of software for data collection, interpreted data and drafted and finalized the manuscript. MO, PO, PZ and BG drafted and finalized the manuscript. All authors read and approved the final manuscript.

Acknowledgement

MEVIS Medical Solutions provided the software used for the airway examiner.

Funding

The study was part of the EU-grant HEALTH F2-2007-201379 COPACETIC.

Author details

¹University of Groningen, Department of Pulmonary Diseases, University Medical Center Groningen, GRIAC research institute, Groningen, the Netherlands. ²University of Groningen, Department of Epidemiology, University Medical Center Groningen, Groningen, the Netherlands. ³University of Groningen, Department of Radiology, University Medical Center Groningen, Groningen, the Netherlands. ⁴Division of Heart & Lungs, University Medical Center Utrecht, Utrecht, the Netherlands. ⁵Department of Radiology, Radboud University Nijmegen Medical Centre, Nijmegen, the Netherlands. ⁶Fraunhofer MEVIS, Institute for Medical Image Computing, Bremen, Germany.

Received: 24 October 2012 Accepted: 17 January 2013

Published: 28 January 2013

References

1. Takasugi JE, Godwin JD: **Radiology of chronic obstructive pulmonary disease.** *Radiol Clin North Am* 1998, **36**:29–55.
2. Schmidt M, Kuhnigk JM, Krass S, Owsijewitsch M, de Hoop B, Peitgen HO: **Reproducibility of airway wall thickness measurements.** In *Proceedings of the SPIE - The International Society for Optical Engineering*, Volume 7624; 2010:762410–762419.
3. Kurashima K, Hoshi T, Takayanagi N, Takaku Y, Kagiya N, Ohta C, Fujimura M, Sugita Y: **Airway dimensions and pulmonary function in chronic obstructive pulmonary disease and bronchial asthma.** *Respiology* 2011, **17**:79–86.
4. Matsumoto H, Niimi A, Tabuena RP, Takemura M, Ueda T, Yamaguchi M, Matsuoka H, Jinnai M, Chin K, Mishima M: **Airway wall thickening in patients with cough variant asthma and nonasthmatic chronic cough.** *Chest* 2007, **131**:1042–1049.
5. Grydeland TB, Dirksen A, Coxson HO, Pillai SG, Sharma S, Eide GE, Gulsvik A, Bakke PS: **Quantitative computed tomography: emphysema and airway wall thickness by sex, age and smoking.** *Eur Respir J* 2009, **34**:858–865.
6. Kim WJ, Silverman EK, Hoffman E, Criner GJ, Mosenifar Z, Sciruba FC, Make BJ, Carey V, Estepar RS, Diaz A, Reilly JJ, Martinez FJ, Washko GR, NETT Research Group: **CT metrics of airway disease and emphysema in severe COPD.** *Chest* 2009, **136**:396–404.
7. Mair G, Maclay J, Miller JJ, McAllister D, Connell M, Murchison JT, Macnee W: **Airway dimensions in COPD: Relationships with clinical variables.** *Respir Med* 2010, **104**:536–542.
8. Pauwels RA, Buist AS, Calverley PM, Jenkins CR, Hurd SS, GOLD Scientific Committee: **Global strategy for the diagnosis, management, and prevention of chronic obstructive pulmonary disease.** NHLBI/WHO Global Initiative for Chronic Obstructive Lung Disease (GOLD) Workshop summary. *Am J Respir Crit Care Med* 2001, **163**:1256–1276.
9. Nakano Y, Muro S, Sakai H, Hirai T, Chin K, Tsukino M, Nishimura K, Itoh H, Pare PD, Hogg JC, Mishima M: **Computed tomographic measurements of airway dimensions and emphysema in smokers. Correlation with lung function.** *Am J Respir Crit Care Med* 2000, **162**:1102–1108.
10. Matsuoka S, Kurihara Y, Yagihashi K, Hoshino M, Nakajima Y: **Airway dimensions at inspiratory and expiratory multisection CT in chronic obstructive pulmonary disease: correlation with airflow limitation.** *Radiology* 2008, **248**:1042–1049.
11. Nishimura M: **Application of three-dimensional airway algorithms in a clinical study.** *Proc Am Thorac Soc* 2008, **5**:910–914.
12. Hasegawa M, Nasuhara Y, Onodera Y, Makita H, Nagai K, Fuke S, Ito Y, Betsuyaku T, Nishimura M: **Airflow limitation and airway dimensions in chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2006, **173**:1309–1315.

13. Patel BD, Coxson HO, Pillai SG, Agustí AG, Calverley PM, Donner CF, Make BJ, Müller NL, Rennard SI, Vestbo J, Wouters EF, Hiorns MP, Nakano Y, Camp PG, Nasute Fauerbach PV, Srean NJ, Campbell EJ, Anderson WH, Paré PD, Levy RD, Lake SL, Silverman EK, Lomas DA: **Airway wall thickening and emphysema show independent familial aggregation in chronic obstructive pulmonary disease.** *Am J Respir Crit Care Med* 2008, **178**:500–505.
14. Lee YK, Oh YM, Lee JH, Kim EK, Lee JH, Kim N, Seo JB, Lee SD: **Quantitative assessment of emphysema, air trapping, and airway thickening on computed tomography.** *Lung* 2008, **186**:157–165.
15. Martinez CH, Chen YH, Westgate PM, Liu LX, Murray S, Curtis JL, Make BJ, Kazerooni EA, Lynch DA, Marchetti N, Washko GR, Martinez FJ, Han MK, COPDGene Investigators: **Relationship between quantitative CT metrics and health status and BODE in chronic obstructive pulmonary disease.** *Thorax* 2012, **67**:399–406.
16. Mets OM, de Jong PA, van Ginneken B, Gietema HA, Lammers JW: **Quantitative computed tomography in COPD: possibilities and limitations.** *Lung* 2012, **190**:133–145.
17. Yamashiro T, Matsuoka S, Estepar RS, Dransfield MT, Diaz A, Reilly JJ, Patz S, Murayama S, Silverman EK, Hatabu H, Washko GR: **Quantitative assessment of bronchial wall attenuation with thin-section CT: An indicator of airflow limitation in chronic obstructive pulmonary disease.** *AJR Am J Roentgenol* 2010, **195**:363–369.
18. Washko GR, Dransfield MT, Estepar RS, Diaz A, Matsuoka S, Yamashiro T, Hatabu H, Silverman EK, Bailey WC, Reilly JJ: **Airway wall attenuation: a biomarker of airway disease in subjects with COPD.** *J Appl Physiol* 2009, **107**:185–191.
19. Bon JM, Leader JK, Weissfeld JL, Coxson HO, Zheng B, Branch RA, Kondragunta V, Lee JS, Zhang Y, Choi AM, Lokshin AE, Kaminski N, Gur D, Sciruba FC: **The influence of radiographic phenotype and smoking status on peripheral blood biomarker patterns in chronic obstructive pulmonary disease.** *PLoS One* 2009, **4**:e6865.
20. Achenbach T, Weinheimer O, Biedermann A, Schmitt S, Freudenstein D, Goutham E, Kunz RP, Buhl R, Dueber C, Heussel CP: **MDCT assessment of airway wall thickness in COPD patients using a new method: correlations with pulmonary function tests.** *Eur Radiol* 2008, **18**:2731–2738.
21. Orlandi I, Moroni C, Camiciottoli G, Bartolucci M, Pistolesi M, Villari N, Mascalchi M: **Chronic obstructive pulmonary disease: thin-section CT measurement of airway wall thickness and lung attenuation.** *Radiology* 2005, **234**:604–610.
22. Grydeland TB, Dirksen A, Coxson HO, Eagan TM, Thorsen E, Pillai SG, Sharma S, Eide GE, Gulsvik A, Bakke PS: **Quantitative CT measures of emphysema and airway wall thickness Are related to respiratory symptoms.** *Am J Respir Crit Care Med* 2009, **34**:858–865.
23. van Klaveren RJ, Oudkerk M, Prokop M, Scholten ET, Nackaerts K, Vernhout R, van Iersel CA, van den Bergh KA, van 't Westeinde S, van der Aalst C, Thunnissen E, Xu DM, Wang Y, Zhao Y, Gietema HA, de Hoop BJ, Groen HJ, de Bock GH, van Ooijen P, Weenink C, Verschakelen J, Lammers JW, Timens W, Willebrand D, Vink A, Mali W, de Koning HJ: **Management of lung nodules detected by volume CT scanning.** *N Engl J Med* 2009, **361**:2221–2229.
24. Quanjer PH, Tammeling GJ, Cotes JE, Pedersen OF, Peslin R, Yernault JC: **Lung volumes and forced ventilatory flows. Report working party standardization of lung function tests, European community for steel and coal. Official statement of the European respiratory society.** *Eur Respir J Suppl* 1993, **16**:5–40.
25. Weinheimer O, Achenbach T, Bletz C, Duber C, Kauczor HU, Heussel CP: **About objective 3-d analysis of airway geometry in computerized tomography.** *IEEE Trans Med Imaging* 2008, **27**:64–74.
26. Gietema HA, Schilham AM, van Ginneken B, van Klaveren RJ, Lammers JW, Prokop M: **Monitoring of smoking-induced emphysema with CT in a lung cancer screening setting: detection of real increase in extent of emphysema.** *Radiology* 2007, **244**:890–897.
27. Parr DG, Sevenoaks M, Deng C, Stoel BC, Stockley RA: **Detection of emphysema progression in alpha 1-antitrypsin deficiency using CT densitometry; methodological advances.** *Respir Res* 2008, **9**:21.
28. Boyden EA: **A synthesis of the prevailing patterns of the bronchopulmonary segments in the light of their variations.** *Dis Chest* 1949, **15**:657–668.
29. Bland JM, Altman DG: **Statistical methods for assessing agreement between two methods of clinical measurement.** *Lancet* 1986, **1**:307–310.
30. Kim S, Choi IS, Kim YJ, Kim CS, Han ER, Park DJ, Kim DE: **Airway responsiveness to inhaled aspirin is influenced by airway hyperresponsiveness in asthmatic patients.** *Korean J Intern Med* 2010, **25**:309–316.
31. Venegas JG, Winkler T, Musch G, Vidal Melo MF, Layfield D, Tgavalekos N, Fischman AJ, Callahan RJ, Bellani G, Harris RS: **Self-organized patchiness in asthma as a prelude to catastrophic shifts.** *Nature* 2005, **434**:777–782.
32. Hogg JC, Chu F, Utokaparch S, Woods R, Elliott WM, Buzatu L, Cherniack RM, Rogers RM, Sciurba FC, Coxson HO, Pare PD: **The nature of small-airway obstruction in chronic obstructive pulmonary disease.** *N Engl J Med* 2004, **350**:2645–2653.
33. Ong KC, Earnest A, Lu SJ: **A multidimensional grading system (BODE index) as predictor of hospitalization for COPD.** *Chest* 2005, **128**:3810–3816.
34. Jones PW, Quirk FH, Baveystock CM, Littlejohns P: **A self-complete measure of health status for chronic airflow limitation. The St. George's Respiratory Questionnaire.** *Am Rev Respir Dis* 1992, **145**:1321–1327.
35. Camiciottoli G, Bigazzi F, Bartolucci M, Cestelli L, Paoletti M, Diciotti S, Cavigli E, Magni C, Buonasera L, Mascalchi M, Pistolesi M: **BODE-index, modified BODE-index and ADO-score in chronic obstructive pulmonary disease: relationship with COPD phenotypes and CT lung density changes.** *COPD* 2012, **9**:297–304.
36. Mets OM, Buckens CF, Zanen P, Isgum I, van Ginneken B, Prokop M, Gietema HA, Lammers JW, Vliegenthart R, Oudkerk M, van Klaveren RJ, de Koning HJ, Mali WP, de Jong PA: **Identification of chronic obstructive pulmonary disease in lung cancer screening computed tomographic scans.** *JAMA* 2011, **306**:1775–1781.

doi:10.1186/1465-9921-14-11

Cite this article as: Dijkstra et al.: Low-dose CT measurements of airway dimensions and emphysema associated with airflow limitation in heavy smokers: a cross sectional study. *Respiratory Research* 2013 **14**:11.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

