

RESEARCH

Open Access



Airways resistance and specific conductance for the diagnosis of obstructive airways diseases

Marko Topalovic¹, Eric Derom², Christian R. Osadnik^{3,4,5,6}, Thierry Troosters^{1,3}, Marc Decramer¹, Wim Janssens^{1*}
and on behalf of the Belgian Pulmonary Function Study Investigators

Abstract

Background: Airway resistance (R_{AW}) and specific airway conductance (sG_{AW}) are measures that reflect the patency of airways. Little is known of the variability of these measures between different lung diseases. This study investigated the contribution of R_{AW} and sG_{AW} to a diagnosis of obstructive airways disease and their role in differentiating asthma from COPD.

Methods: 976 subjects admitted for the first time to a pulmonary practice in Belgium were included. Clinical diagnoses were based on complete pulmonary function tests and supported by investigations of physicians' discretion. 651 subjects had a final diagnosis of obstructive diseases, 168 had another respiratory disease and 157 subjects had no respiratory disease (healthy controls).

Results: R_{AW} and sG_{AW} were significantly different ($p < 0.0001$) between obstructive and other groups. Abnormal R_{AW} and sG_{AW} were found in 39 % and 18 % of the population, respectively, in which 81 % and 90 % had diagnosed airway obstruction. Multiple regression revealed sG_{AW} to be a significant and independent predictor of an obstructive disorder. To differentiate asthma from COPD, R_{AW} was found to be more relevant and statistically significant. In asthma patients with normal FEV_1/FVC ratio, both R_{AW} and sG_{AW} were more specific than sensitive diagnostic tests in differentiating asthma from healthy subjects.

Conclusions: R_{AW} and sG_{AW} are significant factors that contribute to the diagnosis and differentiation of obstructive airways diseases.

Keywords: Body-plethysmography, Airway resistance, Pulmonary function tests, Chronic obstructive pulmonary disease, Asthma

Background

Whole-body plethysmography is an increasingly frequent and important component of comprehensive pulmonary function testing in respiratory medicine, and an integral diagnostic procedure to evaluate static lung volumes and airway resistance [1, 2]. To determine airways resistance, plethysmography applies the gas law of Boyle-Mariotte [3] to evaluate the difference in the pressure of the closed box (in which the subject is breathing) in conjunction with flows measured at the subject's mouth by breathing out of the box [4, 5].

The two most commonly used resistive parameters are airway resistance (R_{AW}) and specific airway conductance (sG_{AW}). R_{AW} reflects changes in alveolar pressure over changes in flow [6] representing true resistance of the airways. R_{AW} is determined by airway narrowing and is therefore considered a good parameter for the diagnosis of airflow obstruction [6]. As the manoeuvre to determine airway resistance is dependent on thoracic gas volume (with larger volumes resulting in opened airways), specific airway resistance (sR_{AW}) is divided by thoracic gas volume to obtain R_{AW} . Mathematically, sR_{AW} does not represent resistance but the flow-standardized work that one needs to perform in order to complete the manoeuvre [4]. sG_{AW} is the inverse of sR_{AW} and therefore reflects the conductance of the airways independent of lung volumes.

* Correspondence: Wim.Janssens@uzleuven.be

¹Department of Clinical and Experimental Medicine, Laboratory of Respiratory Diseases, University Hospital Leuven, KULEUVEN University of Leuven, Herestraat 49, 3000 Leuven, Belgium

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

Despite a large body of physiological evidence, clinicians have lost their interest in measures of resistance which are generally perceived as useless in the work-up of clinical problems. A strong case for the use of such measures in clinical practice recently emerged from a large multicentre study in Belgium (the Belgian Pulmonary Function Study). This study demonstrated that regular measurement of airway resistance (both R_{AW} and sG_{AW}), one of four most commonly used pulmonary function tests (PFTs), significantly contributes to a reduction in the number of differential diagnoses and accuracy of final diagnosis [7]. Nevertheless, it is not yet clear how such parameters vary between different lung disease groups and the extent to which they may contribute to differentiation between asthma, COPD and healthy subjects. The aims of the study were to a) explore the variability of resistance parameters between established diagnoses of respiratory diseases in a general patient sample, and b) investigate whether measures of resistance can differentiate between the obstructive diseases and further refine the diagnosis of asthma or COPD.

Methods

Study subjects

In total 976 subjects were included from the Belgian Pulmonary Function Study. This prospective cohort study includes all successive new subjects from the periods of June 6th till September 12th 2011, and January 16th till June 12th 2012 across 33 hospitals in Belgium. Details on the protocol can be found in the corresponding publication [7]. All subjects were of Caucasian race aged 18 to 75 years with a history of respiratory complaints (dyspnea, cough, sputum, wheezing etc.). All performed complete PFTs at cohort entry including post-bronchodilator spirometry, measurement of diffusion capacity and body-plethysmography including measures of lung volumes, airway resistance (R_{AW}) and conductance (sG_{AW} , only available in 778 subjects). Pulmonologists performed additional tests such as imaging, ECG and other PFTs to determine diagnosis, where necessary. All final diagnoses for each included subject were validated by local Belgian focus groups of 20 to 25 pulmonologists who jointly evaluated all test outcomes. A final diagnosis was selected from a predefined list of 13 diseases covering the wide spectrum of respiratory diseases that are potentially diagnosed by pulmonary function: asthma, COPD, other obstructive diseases, upper airway obstruction, neuromuscular disease, thoracic or pleural disease, obesity, interstitial lung disease, systemic sclerosis or vascular disease, cardiac failure, hyperventilation, no primary pulmonary abnormality, and others (lung cancer, rhinosinusitis, etc.). The study was registered on clinicaltrials.gov (identifier: NCT01297881).

Pulmonary function tests

All PFTs were performed according to the ATS/ERS criteria [8] using standardized equipment available in the participating pulmonary function laboratory (including: SensorMedics Whole Body Plethysmograph, Care fusion, Belgium; Masterscreen Jaeger, Care fusion, Belgium; Medisoft boybox 5500, Belgium; Ganshoren-Medisoft Hyp'air compact, Belgium). Spirometry data are post-bronchodilator measures and expressed, together with airway resistance and lung volume measurements as percent predicted of normal reference values [9, 10]. sG_{AW} was calculated by implementing a least-squared fit of the line through the specific resistance loop at a defined fixed flow of ± 0.5 L/s [5, 11]. Diffusing capacity (DL_{CO}) was measured by the single-breath carbon monoxide gas transfer method and expressed as percent predicted of reference values [12].

Initial inspection of resistance values (expressed as % predicted) from healthy subjects (no primary pulmonary disease) revealed slightly different resistance values than anticipated. This may be due to limited setting where existing reference values for resistive parameters were developed. We therefore established new upper and lower limits of normality for R_{AW} and sG_{AW} based on a 90 % reference interval from our large healthy reference population ($N = 157$). The new proposed upper limit of normality for R_{AW} was 0.38 (kPa/L/s), corresponding to 173 % of the median healthy population reference value. The new lower limit of normality for sG_{AW} was 0.63 (1/kPa*sec) corresponding to 74 % of the reference value. As with existing reference equations on R_{AW} (median value 0.22) and sG_{AW} (median value 0.85), no gender separation was made as no influence on median resistive values of our healthy reference population was observed.

To explore independency of R_{AW} and sG_{AW} on an individual patient level, we compared it with other lung function parameters indicative of the presence of airflow obstruction. These included: FEV₁/FVC ratio, MMEF, RV, TLC and FRC. If one of the latter was out of the predicted normal range, it was considered to be disturbed. Increases in the number of disturbed measures were associated with increased probability of the presence of 'true' airflow obstruction.

Statistical analysis

Statistical Analysis System (SAS) version 9.3, (SAS Institute, Cary, USA) was used for all statistical analyses. A Shapiro-Wilk test was used to explore data normality of each group, and differences between two groups were analysed via unpaired t-tests or Mann-Whitney tests where normal or non-normal data distribution existed, respectively. Logistic-regression models were applied for analysis of binary variables using stepwise selection to identify the subset of variables with the strongest relation to the

presence of obstruction or specific disease diagnoses. The default criterion for variables entering or egressing the models was significance at the 0.15 level. Receiver operating characteristic (ROC) curve analysis was performed via GraphPad Prism version 5.01, (GraphPad Software, La Jolla, California, USA) to identify the best cut-off values of R_{AW} and sG_{AW} to distinguish between asthma and COPD diagnoses. New reference values were determined using MedCalc 14.8.1 (MedCalc Software bvba, Ostend, Belgium), with Tukey test applied to detect outliers.

Results

Baseline characteristics

Baseline characteristics of all included subjects are summarised in Table 1 according to three groups, based upon final diagnosis. The 'healthy' group included subjects without primary pulmonary abnormality; the 'obstruction' group included subjects with COPD, asthma, upper airway obstruction and other obstructive diseases (e.g. chronic bronchitis, bronchiolitis, small airways disease, bronchiectasis, cystic fibrosis); and the 'other pulmonary diseases' group comprised subjects with other, non-obstructive pulmonary diseases. Consistent with clinical practice, the 'obstruction' group was the most common diagnostic group ($N = 651$, 67 % of total sample), followed by 'other pulmonary diseases' ($N = 168$, 17 %) and 'healthy' ($N = 157$, 16 %).

Value of resistance parameters for diagnosis of obstructive airways disease

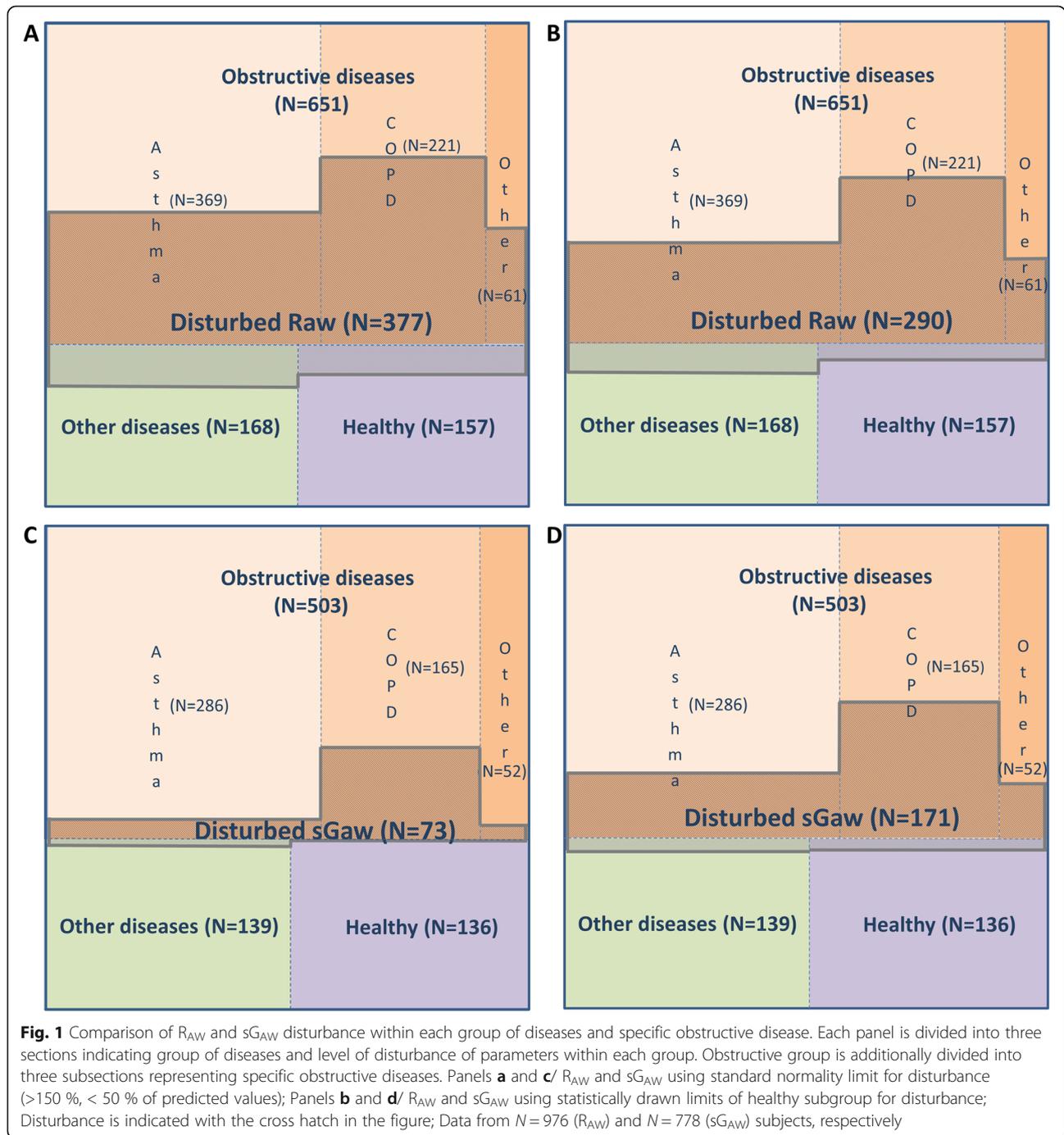
Median R_{AW} values were greater and median sG_{AW} values lower in the group of obstructive patients compared to both the healthy controls and the group of

non-obstructive respiratory diseases (Table 1). These differences were highly statistically significant (both $p < 0.0001$). Expressed as a percentage of predicted values using the reference equation of Quanjer [10], the total study cohort proportion of disturbed R_{AW} (defined as >150 % predicted of the fixed median value) was 39 % ($N = 377$), whilst for sG_{AW} (defined as <50 % predicted) it was 9 % ($N = 73$). Eighteen percent of healthy controls had an abnormal R_{AW} and one had an abnormal sG_{AW} . Implementation of the lower and upper limits of normal for R_{AW} and sG_{AW} developed from our own healthy population resulted in a significant change in the proportion of detected abnormalities. Increased R_{AW} (0.38 kPa/L/s; >173 % predicted) was observed in only 30 % ($N = 290$) of subjects, whilst reduced sG_{AW} (0.63 1/kPa*sec; <74 % predicted) was observed in 22 % ($N = 171$) of subjects (Fig. 1). Independently of the cutoffs used, the group of patients with obstructive diseases had a significant higher proportion of increased R_{AW} . In particular, 47 % of all subjects with obstructive airways disease (Panel A) or 37 % if using the less stringent cut-off (Panel B), presented with an increased resistance. A reduced sG_{AW} (Panels C and D) had a high positive predictive value for obstructive diseases, as 92 % or 90 % (depending of cut-off) of all cases with low sG_{AW} presented with such a diagnosis. This was confirmed in the multiple logistic regression analysis, which identified sG_{AW} as a significant contributing factor to a diagnosis of obstructive lung disease (Table 2). To verify whether resistance parameters (expressed as percent predicted) are beneficial and additional to other PFT parameters to diagnose obstructive airflow disorders, we created different regression models comprising parameters of spirometry only, parameters from spirometry and body-plethysmography and by the combination of the former including diffusing

Table 1 Study population characteristics

	Healthy	Obstruction	Other pulmonary diseases
Subjects, n	157	651	168
Age, years	53 (41–64)	55 (42–65)	57 (44–67)
Smoking, pack yr.	0 (0–5)	0 (0–30)	0 (0–15)
BMI, kg/m ²	25.3 (21.6–28.7)	25.2 (21.1–29.7)	27.6 (23.1–32.0)
FEV ₁ , %predicted	103.3 (91.5–114.7)	84.2 (67.0–97.7)	88.1 (74.4–103.8)
FVC, %predicted	106.6 (97.2–114.9)	97.7 (84.4–110.4)	92.1 (76.5–107.8)
DL _{CO} , %predicted	86.4 (77.0–99.2)	78.7 (63.0–89.9)	70.3 (56.3–85.7)
RV, %predicted	106.9 (91.6–129.7)	131.6 (104.0–159.9)	101.8 (80.9–124.6)
TLC, %predicted	104.6 (95.7–112.1)	106.0 (94.8–117.1)	88.6 (79.7–103.4)
FRC, %predicted	105.1 (93.4–123.1)	119.4 (99.2–143.3)	95.7 (78.0–116.5)
R_{AW} , kPa/L/s	0.24 (0.18–0.30)	0.32 (0.23–0.45)	0.25 (0.17–0.35)
sG_{AW} , 1/kPa*sec	1.21 (0.95–1.57)	0.84 (0.56–1.17)	1.28 (0.90–1.71)

Definition of abbreviations: *BMI* body mass index, *DL_{CO}* carbon monoxide diffusing capacity, *FEV₁* forced expiratory volume in one second, *FRC* functional residual capacity, *FVC* forced vital capacity, *RV* residual volume, *R_{AW}* airway resistance, *sG_{AW}* specific airway conductance, *TLC* total lung capacity, Values are median and IQR



capacity. Stepwise selection excluded R_{AW} in all models, however sG_{AW} contributed significantly to all lung function models.

The presence of abnormal R_{AW} and sG_{AW} values rarely occurred in isolation and was commonly observed in conjunction with disturbances of other PFT parameters indicative of obstruction (Table 3). This was particularly evident for sG_{AW} where only few subjects had normal values of all other PFT parameters. Of the nine cases of isolated sG_{AW} disturbance, two were diagnosed as healthy,

one had neuromuscular disease, one had another obstructive disease, and five had asthma. Additionally, most subjects who presented with isolated abnormally high R_{AW} values had a diagnosis of asthma with normal lung function. Application of the classical cut-offs resulted in detection of 54 subjects with disturbed resistance, of which 28 had asthma, 11 were considered healthy, one had COPD, five presented with other obstructive diseases and nine had various other respiratory diseases. Our more restrictive cut-off threshold identified 29 instances of

Table 2 Relationship between pulmonary function parameters and the presence of obstruction in multiple logistic regression with stepwise selection

Variables	Odds Ratio (95 % Confidence limit)	p value
1.		
sG _{AW} , %predicted	0.996 (0.994–0.998)	0.0003
MEF75, %predicted	0.980 (0.972–0.988)	<0.0001
FEV ₁ /FVC	0.936 (0.912–0.960)	<0.0001
2.		
TLC, %predicted	1.058 (1.043–1.075)	<.0001
sG _{AW} , %predicted	0.997 (0.995–0.999)	0.0031
MMEF, %predicted	0.987 (0.978–0.996)	0.0108
MEF75, %predicted	0.979 (0.969–0.988)	<.0001
FEV ₁ , %predicted	0.970 (0.955–0.986)	<.0001

1/ Variables of spirometry and resistance (initial model1: FVC, %pred, FEV₁, %pred, FEV₁/FVC, MEF75, %pred, MEF50, %pred, MEF25, %pred, MMEF, %pred, PEF, %pred, R_{AW}, %pred, and sG_{AW}, %pred.); 2/ added variables of lung volumes (initial model2: model1, TLC, %pred, RV, %pred, FRC, %pred.); Adding variables of diffusion capacity (initial model3: model2, DL_{CO}, %pred, K_{CO}, %pred.) resulted in same values as model2

disturbed resistance, of which 18 had asthma, six were considered healthy and five had other respiratory diseases.

Value of resistance parameters for differentiating asthma from COPD in obstructive airways diseases

Due to the lack of available data for sG_{AW} in some subjects, diagnostic groups differed in size from the principal analysis, but not in their proportion relative to each other. Median (IQR) R_{AW} values for the asthma (*N* = 369) and COPD (*N* = 221) groups were 0.30 kPa/L/s (0.22–0.42) and 0.39 kPa/L/s (0.26–0.53), respectively (*p* < 0.0001). For sG_{AW} (*N* = 286 for asthma; *N* = 165 for COPD), these were 0.90 1/kPa*sec (0.69–1.22) and 0.60 1/kPa*sec (0.41–0.98), respectively (*p* < 0.0001).

When using the complete set of PFTs in the group of patients with obstructive diseases, application of multiple logarithmic regression with stepwise selection did not reveal an independent statistical role for R_{AW} or sG_{AW} in the differentiation between asthma and COPD. Although R_{AW} was statistically significant in models involving spirometry and plethysmography parameters only, the better variable of differentiation was diffusion capacity (Table 4). When looking at the cumulative disturbances of other PFT parameters per diseased subject, in COPD, the presence of disturbed R_{AW} or sG_{AW} was

Table 4 Relationship between pulmonary function parameters and the presence of asthma in multiple logistic regression with stepwise selection (subjects with obstructive diseases only)

Variables	Odds Ratio (95 % Confidence limit)	p value
1.		
FEV ₁ /FVC	1.043 (1.012–1.074)	<0.0001
MEF25, %predicted	1.024 (1.010–1.038)	0.0024
PEF, %predicted	1.018 (1.006–1.030)	0.0088
R _{AW} , %predicted	1.002 (1.000–1.004)	0.0320
2.		
FEV ₁ , %predicted	1.029 (1.013–1.046)	<.0001
MEF25, %predicted	1.022 (1.010–1.035)	0.0015
R _{AW} , %predicted	1.002 (1.000–1.004)	0.0287
FRC, %predicted	0.981 (0.973–0.989)	<.0001
3.		
DL _{CO} , %predicted	1.071 (1.051–1.090)	<.0001
FEV ₁ /FVC	1.051 (1.024–1.078)	<.0001
TLC, %predicted	1.040 (1.014–1.067)	0.0021
FRC, %predicted	0.970 (0.956–0.984)	0.0041

1/ Variables of spirometry and resistance (initial model1: FVC, %pred, FEV₁, %pred, FEV₁/FVC, MEF75, %pred, MEF50, %pred, MEF25, %pred, MMEF, %pred, PEF, %pred, R_{AW}, %pred, and sG_{AW}, %pred.); 2/ added variables of lung volumes (initial model2: model1, TLC, %pred, RV, %pred, FRC, %pred.); 3/ added variables of diffusion capacity (initial model3: model2, DL_{CO}, %pred, K_{CO}, %pred.)

associated with abnormalities in several other PFT parameters. For those with asthma, increased R_{AW} more commonly occurred in isolation (Table 5). The latter may suggest a more specific role of R_{AW} in the diagnosis of asthma with a non-obstructive spirometry.

Resistance and specific conductance in non-obstructive asthma

In COPD, R_{AW} and sG_{AW} were never independently disturbed as significant changes always concurred with significant changes of other PFT measures, including the FEV₁/FVC ratio by definition. In contrast, subjects with asthma, 28 of 112 (28 %) or 18 of 85 (21 %) presented with isolated R_{AW} disturbance. To further examine this independent value of resistance in asthma, we focussed on the subgroup of asthmatics and healthy controls with normal FEV₁/FVC ratio (above LLN) which resulted in a group of 285 subjects with available R_{AW} data and 213 subjects with available sG_{AW} data. When applying the maximal sum of sensitivity and specificity to select the

Table 3 Disturbed resistance in relation with the disturbance of other lung function parameters indicative of obstruction

R _{AW} (>150 %pred.)		R _{AW} (>173 %pred.)		sG _{AW} (<50 %pred.)		sG _{AW} (<74 %pred.)	
Alone	Together	Alone	Together	Alone	Together	Alone	Together
54 (=20 %)	218	29 (=14 %)	176	1 (=1.5 %)	68	9 (=6 %)	147

Alone: disturbance of only R_{AW} or sG_{AW}. Some patients excluded from analysis due to missing measurement data. Included parameters: FEV₁/FVC, MMEF, RV, TLC and FRC

Table 5 Number of subjects with disturbance per disease (when resistance or specific conductance is already disturbed) stratified over total number of disturbed lung function parameters indicative of obstruction

Disturbance	R_{AW} (>150 %pred.) $N = 204$ (=31 %)		R_{AW} (>173 %pred.) $N = 162$ (=25 %)		sG_{AW} (<50 %pred.) $N = 61$ (=12 %)		sG_{AW} (<74 %pred.) $N = 133$ (=26 %)	
	Asthma ($N = 112$)	COPD ($N = 92$)	Asthma ($N = 85$)	COPD ($N = 77$)	Asthma ($N = 16$)	COPD ($N = 45$)	Asthma ($N = 55$)	COPD ($N = 78$)
0	28	1	18	0	0	0	5	0
1	27	7	23	6	4	0	12	3
2	25	12	19	11	2	2	12	8
3	20	11	16	7	5	4	16	7
4	9	32	6	26	4	19	9	29
5	3	29	3	27	1	20	1	31

Some patients excluded from analysis due to missing measurement data. Proportions observed in a group of all subjects with obstructive diseases. Included parameters: FEV_1/FVC , MMEF, RV, TLC and FRC

best R_{AW} cut-off for asthma diagnosis in that particular subgroup, ROC curve identified an ideal cut-off value of 0.31 kPa/L/s, with a specificity of 76 %, sensitivity of 45 % and positive predictive value of 78 % (Table 6, panel A). For sG_{AW} a cut-off value of 0.98 was identified with associated specificity of 74 %, sensitivity of 50 % and positive predictive value of 75 % (Table 6, panel B).

Discussion

Our study demonstrates measures of airway resistance and specific conductance have a valuable and potentially important role in the diagnosis of obstructive diseases. Specific conductance clearly contributes to diagnosis of obstructive airways diseases, whilst airway resistance can help differentiate between asthma and COPD. The presence of abnormal resistive parameters in isolation from other PFT measures may help differentiate between asthma with normal FEV_1/FVC ratio and normal, good health.

Questions regarding the relevance of resistive parameters for diagnosing respiratory diseases are certainly not new. Prior to this study, it has been shown that specific resistance (reciprocal of specific conductance) may indicate bronchial reversibility in asthma, as it has a different physiological meaning than FEV_1 [13]. Most of its contribution, however, is demonstrated in children with asthma where measurement of specific resistance has been shown to improve diagnostic accuracy

Table 6 Value of resistance to differentiate between healthy subjects and asthma patients with normal FEV_1/FVC ratio

		Asthma (N)	Control (N)	Total (N)
A	$R_{AW} > 0.31$	129	37	166
	$R_{AW} \leq 0.31$	156	120	276
	Total (N)	285	157	442
B	$sG_{AW} < 0.98$	106	35	141
	$sG_{AW} \geq 0.98$	107	101	208
	Total (N)	213	136	349

and decrease diagnostic delay [14–16]. Interestingly, it has also been shown that specific resistance has a better role in predicting asthma than airway resistance [17], and has higher sensitivity than FEV_1 in terms of detecting lung function changes during methacholine challenges [18]. In COPD, both specific conductance and airway resistance are more sensitive for assessing short-acting bronchodilator effects than FEV_1 [18, 19]. Increased airway resistance and decreased specific conductance have also been detected as early features of mild COPD [20, 21]. Detailed analysis of computer tomography scans shows that increased airway resistance in COPD is correlated with the thickening of the airway walls [22]. Our data are in line with these earlier observations, but provide a more comprehensive overview of the role of resistance and specific conductance in a large unbiased sample of a routine respiratory practice in conjunction with other pulmonary function tests. Although resistance and specific conductance overlap with the disturbed pattern of pulmonary function tests in patients with a respiratory disorder, in some subtypes, they may reveal unique components to suggest a certain diagnosis. We demonstrate that both parameters have their specific characteristics which are not the leading but yet still significant contributors to respiratory disease diagnosis.

As expected, the highest frequency of abnormal resistance and specific conductance was found in patients with obstructive airways diseases, particularly COPD. R_{AW} abnormalities were present in 50 % of subjects with mild COPD stages (former GOLD I and II, 75 % of all COPD cases), This proportion rose to 91 % in those with more severe disease (former GOLD III and IV). Whilst fewer subjects had disturbed sG_{AW} , specific conductance appeared more useful than resistance in distinguishing obstructive diseases from non-obstructive, healthy or restrictive diseases. To some extent this may be explained by the fact that sG_{AW} is independent of the thoracic gas volume [23] and may thus differentiate disturbed resistances of smaller airways at low lung volumes (e.g. interstitial lung

disease) from real intraluminal narrowing or collapse, in cases of asthma and COPD [24]. The observation that R_{AW} pops up again when it comes to the further differentiation of asthma from COPD, may be explained by the fact that disproportionately low resistances (by high thoracic gas volumes) for the level of airflow limitation and FEV_1 reduction, is a specific characteristic of COPD [25]. Moreover, our data demonstrate that an isolated increase of R_{AW} might be associated with a diagnosis of asthma when airflow limitation on spirometry is not present. Mechanistically, disturbed FEV_1 measures may be seen as a consequence of increased airway resistance from central and peripheral airways. However, FEV_1 is obtained from the first second of forced expiration and therefore poorly representative of the small airways. The latter are better visualized at the end of a forced expiration or eventually, by direct measures of resistance during tidal breathing [26, 27].

The most appropriate predictive cut-off values to identify normality of resistive and conductive parameters are contentious. A thorough evaluation of our healthy subject population using the current recommended reference equations yielded unsatisfactory results and indicated that the current reference values may not be up to date [10, 28]. Evaluation of these data using the predicted median and 90 % confident intervals in our healthy reference population (which was considered healthy after an extensive evaluation including all necessary tests) showed the cut-offs for R_{AW} to be too liberal and the cut-offs for sG_{AW} too restrictive. Despite calls for revision of these reference equations from different authors, no official action has been taken yet [28, 29]. Solving this problem may lead to a more accurate and better usability of this test in clinical settings as current predictive values were not constituted in a large and healthy representative population.

An inherent limitation of our multi-centre study may lie in its design to measure pulmonary function using variety of equipment, since it can affect variability and consistency of obtained measurements. Weaknesses of airway resistance and specific conductance are also apparent within the analysis. For example, it is clear that both resistive parameters have large variability within core groups, making them weak independent factors for disease differentiation [30]. Another weakness lies in the estimation process for specific conductance, which affects estimation of airway resistance as well. Practically, the line fitted through the resistance loop to calculate sG_{AW} does not always accurately reflect the entire loop pattern. Often, these differences can be visually detected (e.g. widening of the loop, club-shaped format, S pattern, etc. [4]) but may result in similar inclinations. Additional exploration specifically focused on the pattern of such resistive loops will increase the value of these

parameters and may result in a better description of certain phenotypes.

Taken together, airway resistance and specific conductance are certainly not the grand slam winning parameters of pulmonary function tests, however they can significantly contribute to the diagnosis of obstructive diseases and further disease differentiation. Our data also indicate that changing the upper limit of normality for R_{AW} to 0.38 kPa/L/s and accepting a new lower limit of normality for sG_{AW} (0.63 1/kPa*sec) may even improve their diagnostic value.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

Study concept and design: MT, ED, MD, WJ; Analysis and interpretation of the data: MT, TT, WJ; Writing of the manuscript and critical revision: All authors. All authors read and approved the final manuscript.

Acknowledgements

We thank all the pulmonologists, the pulmonary function technicians, the patients and the hospitals who participated in the study for providing data. We thank the Belgian Society of Pneumology for their financial and moral support. Finally, we thank Mrs. Kristien De Bent, Ms. Erica Balligand and Mr. Geert Celis for the provided technical support.

The Belgian Pulmonary Function study investigators

R.Vanherreweghe (Algemeen Stedelijk Ziekenhuis, Aalst); R.Deman, S.Deryke, B.Ghesyens, M.Haerens, S.Maddens (AZ Groeninge, Kortrijk); W.Bultynck, W.Temmerman, L.Van Zandweghe (AZ Sint Blasius, Dendermonde); R.De Pauw, C.Depuydt, C.Haenebalcke, V.Ringoet, D. Van Renterghem (AZ Sint Jan, Brugge); A.Carlier, D.Colle, C.De Cock, J.Lamont (AZ Maria Middelaers, Gent); P.Brancaleone (CH Jolimont-Lobbes); M.Vander Stappen (CHR Haute Senne); R.Peché, Ph.Pierard, P.Quarré, A.Van Meerhaeghe (CHU, Charleroi); D.Cataldo, J.L.Corhay, B.Duysinx, V.Heinen, M.Naldi, D.Nguyen, L.Renaud (CHU, Liège); M.Cotils, P.Duchatelet, A.Gocmen, C.Lenclud (Clinique Louis Caty-Baudour); Ph.Lebrun, J.Noel (Clinique Saint-Pierre, Ottignies); A.Frémault (Grand Hôpital de Charleroi); Ph.Bertrand, B.Bouckaert, I.Demedts, P.Demuyne (Heilig Hart Ziekenhuis, Roeselare); E.Frans, A.Heremans, T.Lauwerier, J.Roelands (Imelda, Bonheiden); G.Bral, I.Declercq, I.Malysse (Jan Ypermanziekenhuis, Ieper); V.Van Damme (St. Andries Ziekenhuis, Tielt); Dr.Martinot (St. Elisabeth, Namur); P.Vandenbrande (St. Maarten Duffel, Mechelen); M.Bruijnneel, I.Muylle, V.Ninane, (St. Pierre, Bruxelles); Ph.Collard, G.Liistro, B.Mwenge Gimbada, Th.Pieters, C.Pilette, F.Pirson, D.Rodenstein (UCL, Bruxelles); L.Delaunois, E.Marchand, O.Vandenplas (UCL, Mont-Godinne); W.De Backer, P.Germonpre, A.Janssens, A.Vrints (UZA, Antwerpen); M.Decramer, W.Janssens, P.Van Bleyenbergh, G.Verleden, W.Wuyts (UZ Gasthuisberg, Leuven); G.Brusselle, E.Derom, G. Joos, K.Tournoy, K.Vermaelen (UZ, Gent); P.Alexander, C.Gysbrechts (Ziekenhuis Ronse); L.Bedert, Dr.Bomans, Dr.De Beukelaar, E.De Droogh, D.Galdermans, Dr.Engelbrecht, Dr.Lefebure, H.Slabbynck, Dr.Van Mulders, Dr.Van Schaardenburg, Dr.Verbuyst (ZNA, Antwerpen); M.Daenen (ZOL Genk).

Funding

Astra Zeneca Chair 2012–2014. Belgian Society of Pneumology. WJ is supported by the Flemish Research Foundation (FWO).

Author details

¹Department of Clinical and Experimental Medicine, Laboratory of Respiratory Diseases, University Hospital Leuven, KULEUVEN University of Leuven, Herestraat 49, 3000 Leuven, Belgium. ²Respiratory Division, University Hospital Ghent, University of Ghent, Ghent, Belgium. ³Department of Rehabilitation Sciences, Faculty of Kinesiology and Rehabilitation Sciences, KU Leuven, Leuven, Belgium. ⁴Department of Physiotherapy, Monash University, Victoria, Australia. ⁵Institute for Breathing and Sleep, Victoria, Australia. ⁶Monash Health, Monash Lung and Sleep, Victoria, Australia.

Received: 18 March 2015 Accepted: 8 July 2015

Published online: 22 July 2015

References

- Coates AL, Peslin R, Rodenstein D, Stocks J. Measurement of lung volumes by plethysmography. *Eur Respir J*. 1997;10:1415–27.
- Pellegrino R, Viegi G, Brusasco V, Crapo RO, Burgos F, Casaburi R, et al. Interpretative strategies for lung function tests. *Eur Respir J*. 2005;26:948–68.
- Dubois AB, Botelho SY, Comroe Jr JH. A new method for measuring airway resistance in man using a body plethysmograph: values in normal subjects and in patients with respiratory disease. *J Clin Invest*. 1956;35:327–35.
- Crieie CP, Soricther S, Smith HJ, Kardos P, Merget R, Heise D, et al. Body plethysmography—its principles and clinical use. *Respir Med*. 2011;105:959–71.
- Stocks J, Godfrey S, Beardsmore C, Bar-Yishay E, Castile R. Plethysmographic measurements of lung volume and airway resistance. ERS/ATS Task Force on Standards for Infant Respiratory Function Testing. European Respiratory Society/ American Thoracic Society. *Eur Respir J*. 2001;17:302–12.
- Shiner RJ, Steier J. Lung Function Tests Made Easy. UK: Elsevier Health Sciences; 2012.
- Decramer M, Janssens W, Derom E, Joos G, Ninane V, Deman R, et al. Contribution of four common pulmonary function tests to diagnosis of patients with respiratory symptoms: a prospective cohort study. *Lancet Respir Med*. 2013;1:705–13.
- Miller MR, Hankinson J, Brusasco V, Burgos F, Casaburi R, Coates A, et al. Standardisation of spirometry. *Eur Respir J*. 2005;26:319–38.
- Quanjer PH, Tammeling GJ, Cotes JE, Pedersen OF, Peslin R, Yernault JC. [Lung volumes and forced ventilatory flows. Work Group on Standardization of Respiratory Function Tests. European Community for Coal and Steel. Official position of the European Respiratory Society]. *Rev Mal Respir*. 1994;11 Suppl 3:5–40.
- Quanjer P. Standardized lung function testing. *Bull Europ Physiopath Resp*. 1983;19:33–8.
- Hantos Z, Galgoczy G, Daroczy B, Dombos K. Computation of the equivalent airway resistance: a comparison with routine evaluations of plethysmographic measurements. *Respiration*. 1978;36:64–72.
- Miller A, Thornton JC, Warshaw R, Anderson H, Teirstein AS, Selikoff IJ. Single breath diffusing capacity in a representative sample of the population of Michigan, a large industrial state. Predicted values, lower limits of normal, and frequencies of abnormality by smoking history. *Am Rev Respir Dis*. 1983;127:270–7.
- Mahut B, Peiffer C, Bokov P, Delclaux C, Beydon N. Use of specific airway resistance to assess bronchodilator response in children. *Respirology*. 2011;16:666–71.
- Brzozowska A, Majak P, Grzelewski T, Stelmach W, Kaczmarek J, Stelmach P, et al. Measurement of specific airway resistance decreased the risk of delay in asthma diagnosis in children. *Allergy Asthma Proc*. 2009;30:47–54.
- Kirkby J, Stanojevic S, Welsh L, Lum S, Badier M, Beardsmore C, et al. Reference equations for specific airway resistance in children: the Asthma UK initiative. *Eur Respir J*. 2010;36:622–9.
- Nielsen KG, Bisgaard H. Discriminative capacity of bronchodilator response measured with three different lung function techniques in asthmatic and healthy children aged 2 to 5 years. *Am J Respir Crit Care Med*. 2001;164:554–9.
- Buhr W, Jorres R, Knapp M, Berdel D. Diagnostic value of body plethysmographic parameters in healthy and asthmatic young children is not influenced by breathing frequency. *Pediatr Pulmonol*. 1990;8:23–8.
- Van Noord JA, Smeets J, Clement J, Van de Woestijne KP, Demedts M. Assessment of reversibility of airflow obstruction. *Am J Respir Crit Care Med*. 1994;150:551–4.
- Borrill ZL, Houghton CM, Woodcock AA, Vestbo J, Singh D. Measuring bronchodilation in COPD clinical trials. *Br J Clin Pharmacol*. 2005;59:379–84.
- Deesomchok A, Webb KA, Forkert L, Lam YM, Ofir D, Jensen D, et al. Lung hyperinflation and its reversibility in patients with airway obstruction of varying severity. *COPD*. 2010;7:428–37.
- Dias RM, Chacur FH, Carvalho SR, Neves DD. Which functional parameters can help differentiate severe asthma from COPD? *Rev Port Pneumol*. 2010;16:253–72.
- Kosciuch J, Krenke R, Gorska K, Zukowska M, Maskey-Warzechowska M, Chazan R. Relationship between airway wall thickness assessed by high-resolution computed tomography and lung function in patients with asthma and chronic obstructive pulmonary disease. *J Physiol Pharmacol*. 2009;60 Suppl 5:71–6.
- Briscoe WA, Dubois AB. The relationship between airway resistance, airway conductance and lung volume in subjects of different age and body size. *J Clin Invest*. 1958;37:1279–85.
- Gibson GJ. *Clinical Tests of Respiratory Function* 3rd Edition. Taylor & Francis; 2008
- Kaminsky DA. What does airway resistance tell us about lung function? *Respir Care*. 2012;57:85–96.
- Bassiri AG, Girgis RE, Doyle RL, Theodore J. Detection of small airway dysfunction using specific airway conductance. *Chest*. 1997;111:1533–5.
- O'Donnell DE, Laveneziana P, Ora J, Webb KA, Lam YM, Ofir D. Evaluation of acute bronchodilator reversibility in patients with symptoms of GOLD stage I COPD. *Thorax*. 2009;64:216–23.
- Koch B, Friedrich N, Volzke H, Jorres RA, Felix SB, Ewert R, et al. Static lung volumes and airway resistance reference values in healthy adults. *Respirology*. 2013;18:170–8.
- Garcia-Rio F, Dorgham A, Pino JM, Villasante C, Garcia-Quero C, Alvarez-Sala R. Lung volume reference values for women and men 65 to 85 years of age. *Am J Respir Crit Care Med*. 2009;180:1083–91.
- Ruppel GL, Enright PL. Pulmonary function testing. *Respir Care*. 2012;57:165–75.

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

