

RESEARCH ARTICLE

A multi-scale model of gas transport in the lung to study heterogeneous lung ventilation during the multiple-breath washout test

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Citation: Hasler D, Anagnostopoulou P, Nyilas S, Latzin P, Schittny J, Obrist D (2019) A multi-scale model of gas transport in the lung to study heterogeneous lung ventilation during the multiple-breath washout test. *PLoS Comput Biol* 15(6): e1007079. <https://doi.org/10.1371/journal.pcbi.1007079>

Editor: Alison L. Marsden, Stanford University, UNITED STATES

Received: November 16, 2018

Accepted: May 1, 2019

Published: June 17, 2019

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Data Availability Statement: All simulation data are available under DOI: [10.5281/zenodo.1476557](https://doi.org/10.5281/zenodo.1476557)

Funding: JS received financial support from the Swiss National Science Foundation (grants #310030_153468 and #310030_175953) <http://www.snf.ch>. DO and PL received financial support from the Commission for Technology and Innovation (CTI) – Swiss Innovation Agency (grants 14435.1 PFLS-LS and 18274.1 PFLS-LS) <https://www.innosuisse.ch/inno/en/home.html>. The

Abstract

The multiple-breath washout (MBW) is a lung function test that measures the degree of ventilation inhomogeneity (VI). The test is used to identify small airway impairment in patients with lung diseases like cystic fibrosis. However, the physical and physiological factors that influence the test outcomes and differentiate health from disease are not well understood. Computational models have been used to better understand the interaction between anatomical structure and physiological properties of the lung, but none of them has dealt in depth with the tracer gas washout test in a whole. Thus, our aim was to create a lung model that simulates the entire MBW and investigate the role of lung morphology and tissue mechanics on the tracer gas washout procedure. To this end, we developed a multi-scale lung model to simulate the inert gas transport in airways of all size. We then applied systematically different modifications to geometrical and mechanical properties of the lung model (compliance, residual airway volume and flow resistance) which have been associated with VI. The modifications were applied to distinct parts of the model, and their effects on the gas distribution within the lung and on the gas concentration profile were assessed. We found that variability in compliance and residual volume of the airways, as well as the spatial distribution of this variability in the lung had a direct influence on gas distribution among airways and on the MBW pattern (washout duration, characteristic concentration profile during each expiration), while the effects of variable flow resistance were negligible. Based on these findings, it is possible to classify different types of inhomogeneities in the lung and relate them to specific features of the MBW pattern, which builds the basis for a more detailed association of lung function and structure.

funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing interests: The authors have declared that no competing interests exist.

Author summary

Obstructive lung diseases, like cystic fibrosis or primary ciliary dyskinesia, lead to inhomogeneous ventilation. The degree of observed inhomogeneity represents a clinical measure for the progression of the disease. The multiple-breath washout (MBW) is a lung function test that measures this inhomogeneity in the lung. However, the factors that influence the results of the test and differentiate between health and disease are not well understood. Computational models help us to understand better the relation between anatomical structure and physiological properties of the lung, but none of them has dealt in depth with the MBW test in whole. Our aim was to create a lung model that simulates the entire MBW test and study the role of lung structure and tissue mechanics on the washout procedure. We developed a multi-scale lung model to simulate the inert gas transport in all airways including the gas exchange area. Our model offers the opportunity to understand the ventilation distribution in the healthy lung. It can also mimic certain patterns of lung disease by applying modifications in mechanical properties out of the physiological limits. Thus, it can be used to study MBW characteristics in health and disease.

Introduction

The multiple-breath washout (MBW) is a lung function test that measures the degree of ventilation inhomogeneity [1] and is increasingly used for both research and clinical purposes in patients with obstructive lung disease, such as cystic fibrosis, primary ciliary dyskinesia, etc. [2–4]. The test is based on the clearance of a tracer gas during multiple tidal breaths. Each MBW test comprises a washin and a washout phase. During the washin phase, the tracer gas (normally an inert extrinsic gas) is delivered in a known concentration. When the tracer gas concentration reaches an equilibrium in the lung, the washout phase starts. In the case of inert intrinsic gases (N_2), the test is simplified, as no washin phase is needed. In the washout phase the subject inhales a gas other than the tracer gas (e.g. pure O_2 in case of N_2 MBW), so that the lungs wash out the tracer gas gradually by each expiration [2]. The progressive decrease in tracer gas concentration during the washout (washout envelope) as well as the breath-by-breath analysis provide useful information about the distribution of ventilation within the lung [1], and for this reason the test is of increasing importance for the medical community. However, the biomechanical phenomena that influence the specific washout profile of a tracer gas are not well understood.

The anatomical diversity in the airways as well as the physiological properties of the lung tissue (e.g. compliance, resistance) influence the respiratory function in a complex way [5, 6]. Over the last years, computational models have been used to better understand those interactions. Anatomically based three-dimensional (3D) lung models have elegantly simulated lung tissue mechanics [7–9] and 3D fluid dynamics in large airways [10–12], but without specific focus on the ventilation and gas transport in the entire lung in the context of MBW. More simplified models have addressed these phenomena, but either have not included physiological asymmetries in lung morphology, or have not modeled the whole lung [13–23]. Therefore, results of these models cannot be directly compared to clinical data in humans [24].

The aim of this article is to introduce a computational model that simulates gas transport in the entire lung during the MBW test, taking into account transport phenomena at different scales. It should allow relating physiological phenomena at the smallest scales of the lung to gas concentrations at the mouth, which can be measured clinically. Such simulations can provide

detailed insight in the gas transport dynamics during the MBW in different airways and help to understand better the effect of lung morphology and tissue mechanics on tracer gas wash-out. To this end, structural data from human lungs have been used to construct a fractal lung model, taking into account morphological and physiological asymmetries in lung anatomy [15, 25, 26] including the number and size distribution of the acini [27, 28]. We have designed the model in a way that is complex enough for the introduction of different types of asymmetries and structural properties of the acini, but simple enough to allow for quick computational turn-around times on desktop computers. The model was used to study some basic mechanisms, which lead to well-known clinical observations in MBW [1], and an example of a model for the healthy lungs was created, using known parameters that can produce a physiological heterogeneous ventilation distribution, as described in healthy individuals [6].

Such models inherently lack suitable validation methods [29] because ground truth data is not available in the lower airways. Here, we compared the MBW simulations in the computational lung model to data from nitrogen MBW (N₂MBW) tests from healthy controls (N = 4). Although this comparison is too small to serve as a statistically solid validation of the model, the good agreement between computational results and in vivo data illustrates the potential of the proposed model to reproduce clinical test data.

Methods

Ethics statement

The study was approved by the Ethics Committee of the Canton of Bern, Switzerland (KEK-Gesuchs-Nr: 181/03), and caregivers gave written informed consent.

Lung morphology

In our model, the airway morphology is represented by a generic dichotomous tree network of straight branching pipes [30], which terminates in trumpet-like compartments. The straight pipes represent the larger, non-compliant airways, where convective transport is dominant. The dimensions of the trachea and the relative dimensions (i.e. with respect to the trachea) of the airways in the first four generations are defined according to anatomical data from Weibel [26]. In addition, the entire airway network of pipes (all conducting airways including the trachea, without the trumpet-like compartments) was then scaled to meet a specific functional residual capacity (FRC) that is, the air volume that remains in the lung after tidal expiration. The length of the scaled trachea was defined as

$$l_0 = l_{0W} \left(\frac{FRC}{FRC_W} \right)^{1/3} \quad (\text{Eq 1})$$

where the subscript W indicates a reference quantity [26].

For airways past the 4th generation, we applied the scheme for a regular branching asymmetry introduced by Majumdar et al. [25, 30]. In this scheme, each pipe-like airway bifurcates in a major and a minor daughter airway. In the airway network every parent, minor, and major daughter share a common node. The dimensions of the daughter airways, namely their diameter and length, are different fractions of the dimensions of their common parent pipe,

$$\begin{aligned} d_{z+1,maj} &= d_z \kappa_{maj} & \text{with } \kappa_{maj} &= (1-r)^{1/\eta} \text{ and} \\ d_{z+1,min} &= d_z \kappa_{min} & \text{with } \kappa_{min} &= r^{1/\eta} \end{aligned} \quad (\text{Eq 2})$$

where d_z is the diameter of a pipe at generation z , and r and η denote the asymmetry parameter

and reduction rate, respectively. The same scheme was used to define the airway lengths. Majumdar et al. proposed values $r = 0.326$ and $\eta = 2.97$ for which the resulting structure best represents the human lung on a statistical basis [25]. In the present model, this bifurcation scheme was applied recursively until the diameter of a pipe was less than a lower limit diameter $d_{lim} = 1.8 \text{ mm}$. This limit defines the terminal pipes in the model and was chosen to represent bronchi at the 8th-12th generation. The generation of the individual terminal pipe-like airways depends on the limit diameter and varies within the model, due to the asymmetric bifurcation scheme. This limit diameter was chosen in order to complete the simulations with reasonable computational costs. A numerical experiment showed very small differences in the simulation outcomes for $d_{lim} = 1.6, 1.8, 2.0 \text{ mm}$.

The computational unit distal to a terminal pipe constitutes a lobule and was modelled using a trumpet-like compartment (trumpet lobule). Please note that the “lobule” as defined here differs from the anatomical term lobule as defined by Miller [31], because it contains, apart from the acini that include respiratory bronchioles, alveolar ducts and alveoli, also the generations of conducting airways with diameter smaller than d_{lim} . Unlike the pipes, the trumpet lobule is a compartment with diverging, time-variable cross-section (Fig 1A). The number of these trumpet lobules is equal to the number of terminal pipes, as each terminal pipe leads to one lobule.

The residual volume of the trumpet lobules was defined such that the total volume of the model (pipes and trumpet lobules) at the end of a tidal expiration equaled a predefined FRC. Fig 1B shows a sketch of a model lung illustrating the relative scales of a network of airways defined by the bifurcation rule (Eq 2).

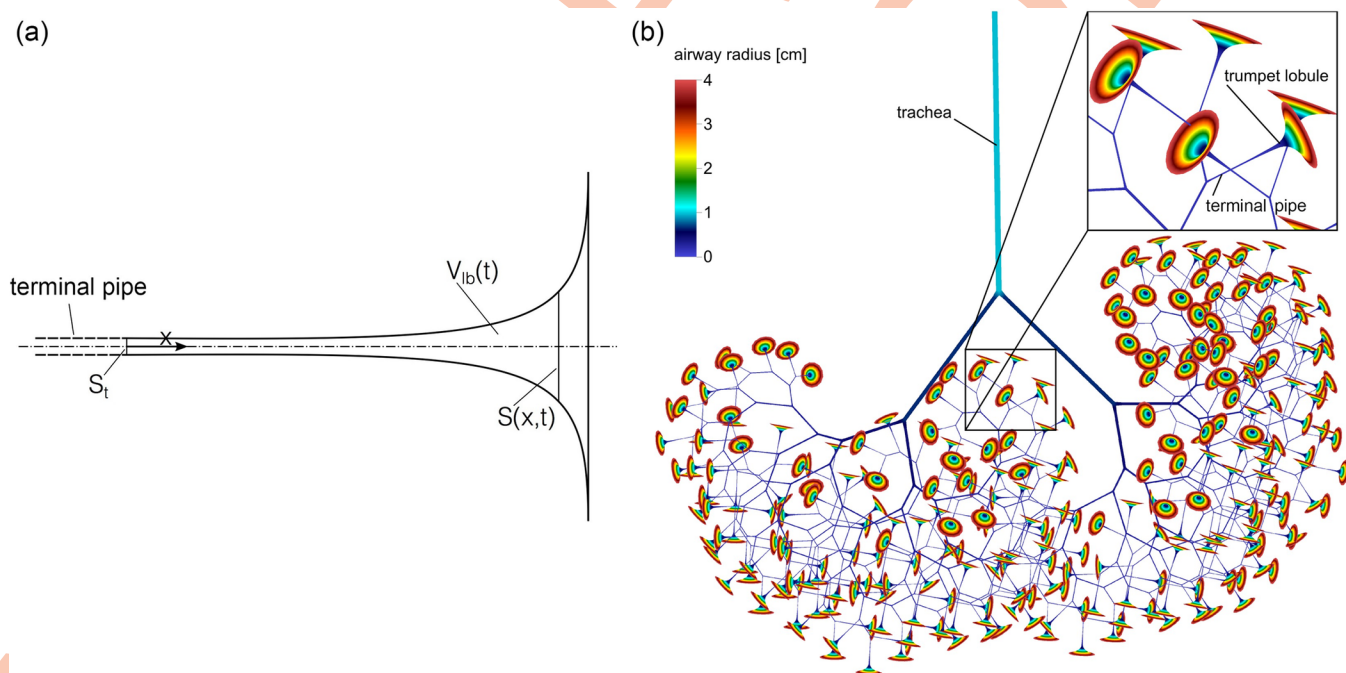


Fig 1. Trumpet lobule geometry and lung model morphology. (a) Schematic representation of the trumpet model used for the lobules with the cross-section of the terminal pipe S_t and the time-variable cross-section $S(x,t)$ and volume $V_{lb}(t)$ of a trumpet lobule model. (b) Idealized lung model morphology based on bifurcation rule (Eq 2). For better visibility of the airway structure, all diameters were scaled with the factor 0.1. The length and diameter ratio of airways until the fourth generation are irregular and follow from empirical data by Weibel [26]. After the fourth generation, airways bifurcate in a minor and major daughter airway according to Eq 2. From this scheme an asymmetric network of airways with different total lengths results. With each generation, airways get smaller, both in terms of length and of cross-section. In the lung model, when the diameter of the airways falls below a limit diameter d_{lim} , a so-called terminal pipe is reached, and different model (the trumpet lobule) is used to represent smaller airways.

<https://doi.org/10.1371/journal.pcbi.1007079.g001>

Mathematical description of the trumpet lobule

For the trumpet model representing the lobules and their peripheral airways, a model for the total cross-section of the trumpet lobule $S_{lb}(x,t)$, as well as for the mean advection velocity $u_{lb}(x,t)$ had to be derived.

The flow rate at the inlet of each trumpet equals the flow rate in a terminal pipe and $Q_t(t)$ was known from the results of a model for lung ventilation (described in the upcoming section) and was used to compute the total volume of the trumpet $V_{lb}(t) = \int_0^t Q_t dt + V_{lb}^0$.

The initial volume of the trumpet lobule, $V_{lb}(t = 0) = V_{lb}^0$, followed from FRC-based scaling of the lung model.

Assuming a uniform homothety ratio of $\kappa = 0.85$ [19] for airways lumped in a trumpet lobule, the total change of cross-section along the streamwise coordinate x can be described as

$$S(x, 0) = S_t \hat{\kappa}^{z(x)}, \text{ with } \hat{\kappa} = 2\kappa^2 \tag{Eq 3}$$

where S_t is the cross-section of the terminal pipe, and $z(x)$ is the generation at position x with respect to the inlet of the trumpet lobule where $z = x = 0$.

Considering l_t to be the length of the terminal pipe, the cumulative length at generation z (with respect to the inlet of the lobule) would be $\sum_{k=1}^z l_t \kappa^k$.

The limits of this sum for $z \rightarrow \infty$ are

$$\lim_{z \rightarrow \infty} \sum_{k=1}^z l_t \kappa^k = l_t \kappa / (\kappa - 1) =: L. \tag{Eq 4}$$

From these relations, an expression for the generation in function of the distance to the inlet of the trumpet can be computed,

$$z(x) = \frac{\log \left[x \frac{\kappa - 1}{\kappa l_t} + 1 \right]}{\log(\kappa)} \tag{Eq 5}$$

Using Eq 3 together with Eq 4 for further treatment of the lobule model becomes a rather cumbersome task. We therefore sought a model $S_{lb}(x,t)$, which approximates Eq 5 but allows to derive an analytical expression for $u_{lb}(x,t)$.

To this end, we used a power law of the form

$$S_{lb}(x, t) = p_1 x^{n_1} + p_2 x^{n_2} + S_t, \tag{Eq 6}$$

with $n_1, n_2 = 20, 2$, where the coefficients p_1 and p_2 were defined such that the lobule cross-section S_{lb} intersects with Eq 3 at a chosen generation z^* , and the prescribed initial volume V_{lb}^0 of the trumpet lobule is obtained for a given length l_{lb} of the trumpet lobule. The mathematical expression for this parameter definition as well as a graphic comparison between the formulation Eq 3 and the model Eq 6 are provided in S1 Appendix (Section 1).

The expression given in Eq 6 determines the shape of the trumpet lobule at any time.

An important feature of this model is the major contribution of peripheral airway (where x is close to l_{lb}) to the overall expansion of the lung. Furthermore, the differentiation with respect to time of Eq 6 as well as the integration along the trumpet centre line coordinate x is straightforward and therefore allows to formulate a modified advection diffusion equation for the trumpet lobule (see S1 Appendix, Section 1)

Apart from the constraint on the total volume, the geometrical and mechanical properties of the stiff (pipes) and the compliant parts (trumpet lobules) can be modified individually to study systematically the effects of structural lung inhomogeneity.

Ventilation

Lung ventilation was simulated by a lumped parameter (0-dimensional) model (Fig 2) based on the model morphology described above. For the pipe-like airways, purely resistive elements were used. Womersley's theory for pulsatile flow in tubes [32] was applied to account for inertial effects during normal breathing, which have a considerable effect on the flow resistance in bigger airways until the fifth generation [33] (see also S1 Appendix). In general, the pressure difference between two subsequent nodes with index i and j in the network of pipe-like airways reads $p_i - p_j = R_{ij}Q_{ij}$. Here Q_{ij} is the flow rate from node i to node j , and the hydrodynamic resistance R_{ij} depends on the radius r_{ij} and the length l_{ij} of the conducting airway between two nodes, and on the breath period T_B . More information on the pressure-flow relation can be found in S1 Appendix.

The trumpet lobule model, mainly representing compliant airways, is composed of a non-linear compliance element (elastic pressure, p_{el}) and a resistance element (viscous pressure loss, p_{diss}), acting in series between a node i , corresponding to a terminal pipe, and the pleural gap with pressure p_{pl} . This representation is based on the pressure-volume relation as presented by Bates [34]. However, instead of the regular linear elastic law, we used an empirical exponential expression to represent p_{el} . The corresponding pressure difference was defined as

$$p_i - p_{pl} = \underbrace{\beta e^{\gamma V_{lb}^0} (e^{\gamma \tilde{V}_{lb}} - 1)}_{p_{el}} + \underbrace{R_{lb} Q_{lb}}_{p_{diss}} \quad (\text{Eq 7})$$

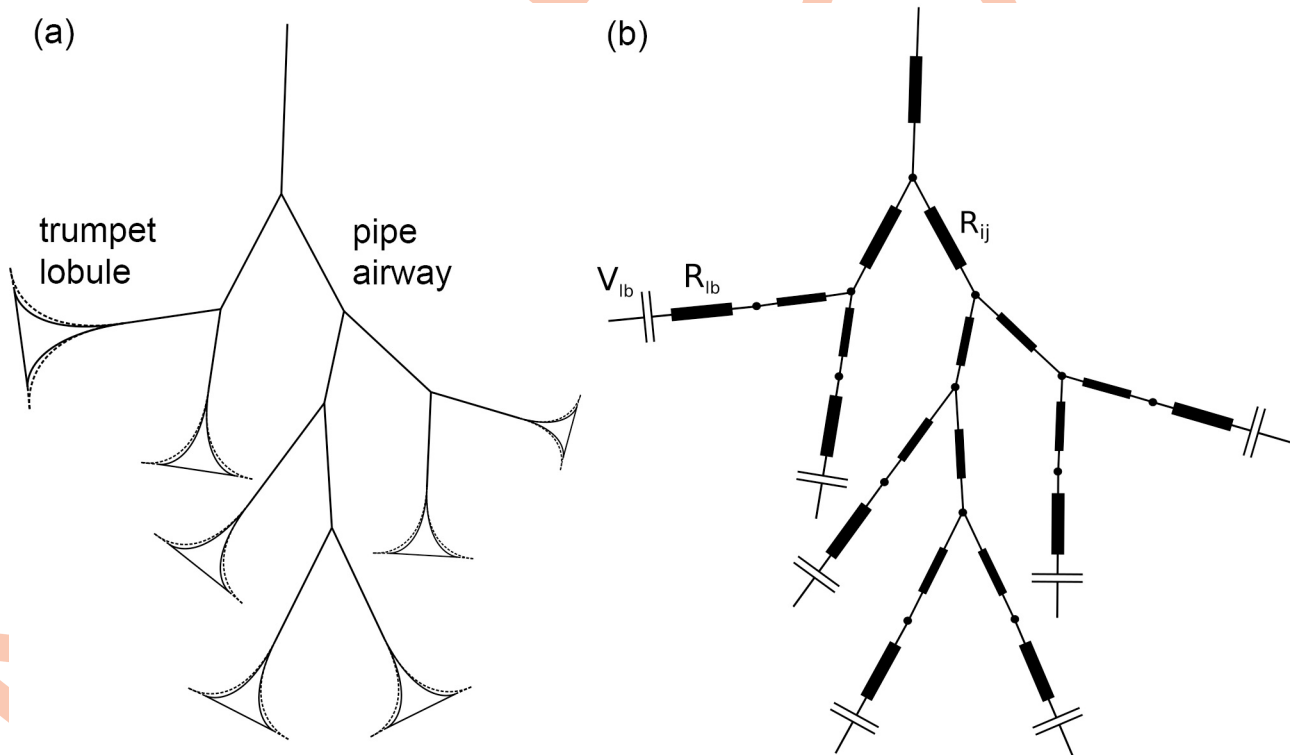


Fig 2. Ventilation lumped parameter model schematics. (a) Network of pipe- and trumpet-like elements used for the upper and lower airways, respectively. (b) Corresponding lumped parameter model composed of resistances for the conducting airways R_{ij} and for the trumpet lobule R_{lb} , and compliance elements which relate the trans-lobular pressure to the volume of the trumpet lobule $V_{lb}(t)$ (see Eq 7).

<https://doi.org/10.1371/journal.pcbi.1007079.g002>

where R_{lb} , $V_{lb}(t) = V_{lb}^0 + \tilde{V}_{lb}(t)$ and Q_{lb} denote the total flow resistance of the lobule, its volume and the flow rate into the trumpet lobule, respectively; V_{lb}^0 is the lobule volume at FRC and $\tilde{V}_{lb}(t)$ is the dynamic volume during breathing [34]. The shape parameters β and γ were used to define and modify the degree of the compliance parameter ϕ (i.e. the distensibility of the lobule, described below in details) and the non-linearity (in the pressure-volume relation) of the trumpet lobule, respectively. There is a single β - γ pair for each lobule, and both are defined indirectly by fixed intersection points in the pressure-volume curve for the elastic pressure (see Fig 3A). More information on the mechanical properties of the model can be found in the Section “Structural and mechanical modifications of the model”.

To ensure mass conservation, the flow rates in and out of each node i satisfy the balance $\Sigma Q = 0$. At the inlet, i.e. upstream node of the trachea, the flow rate is defined by the predefined boundary conditions $Q_{in}(t)$, and for the trumpet lobule the flow rate of the terminal pipe Q_t equals the change of volume of the lobule,

$$Q_t = \frac{dV_{lb}}{dt}. \tag{Eq 8}$$

This results in a system of differential algebraic equations (DAE). Together with appropriate boundary conditions, the DAE system governs the flow distribution in the lumped parameter model. A detailed description of the numerical implementation of the lumped parameter model is provided in S1 Appendix.

During inspiration, gas is transported from the mouth and the nose toward the alveolar membrane, as a result of the motion of the diaphragm and the thoracic cavity, causing a volume increase in the lung and a pressure decrease in the pleural gap. This negative (relative) pressure in the pleural gap causes a pressure gradient across the peripheral lung tissue and along the airways to the mouth and nose. Therefore, the pleural pressure and the pressure at

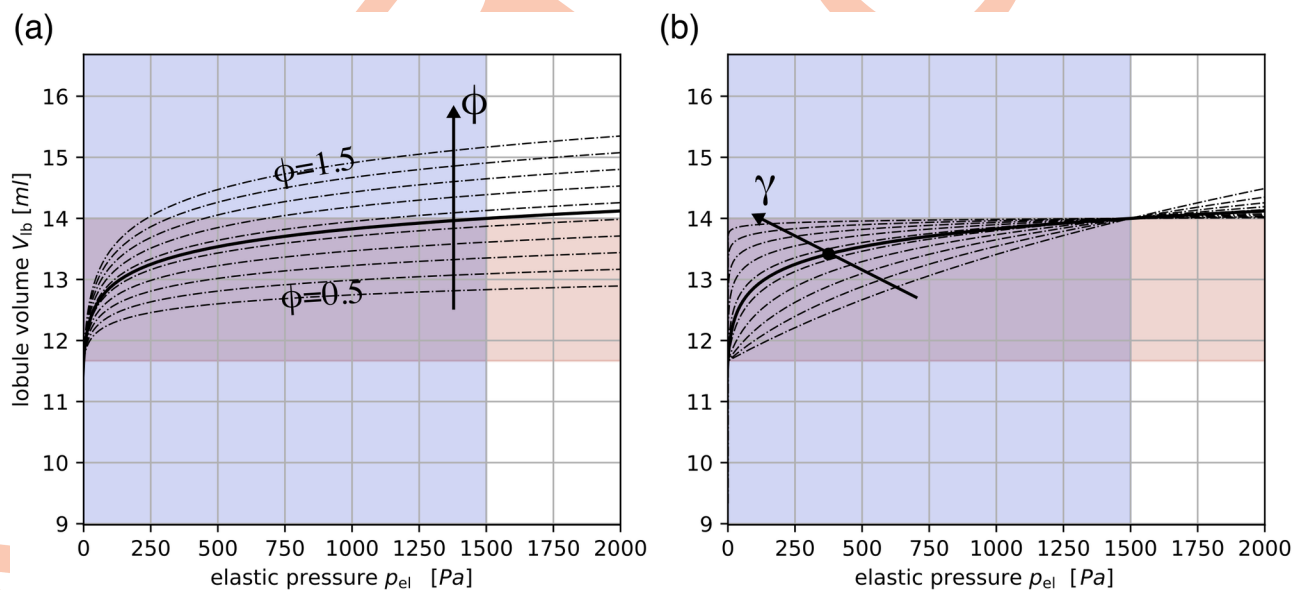


Fig 3. Pressure-volume curve for trumpet lobule. Lobular volume V_{lb} in function of the elastic pressure p_{el} as defined in Eq 3. The purple area indicates the reference pleural pressure range p_{lb}^{TV} and the orange area indicates the nominal lobule volume range $[V_{lb}^0, V_{lb}^0 + V_{lb}^{TV}]$. In panel (a) the influence of the compliance modification parameter $\phi \in [0.5, 1.5]$ is shown. For lobules with normal compliance ($\phi = 1$, solid line) the elastic pressure curve intersects in the point $(p_{lb}^{TV}, V_{lb}^0 + V_{lb}^{TV})$. In panel (b) the effect of the non-linear parameter γ is shown. The parameter was chosen such that the elastic pressure curve intersects with the point $(\frac{1}{4} p_{lb}^{TV}, V_{lb}^0 + \frac{3}{4} \phi V_{lb}^{TV})$ (black dot). Note that $\phi = 1$ was used for all graphs in panel (b).

<https://doi.org/10.1371/journal.pcbi.1007079.g003>

the mouth would be obvious choices for the boundary conditions for the lumped parameter model (LPM). However, the pleural pressure is in general not measurable during clinical routine. Instead, the pleural pressure was determined computationally such that a prescribed flow rate $Q_{in}(t)$ was attained in the trachea. This flow rate could be easily determined during clinical testing.

Gas transport

Gas transport in the lung occurs by convection and diffusion [35, 36]. To model the distribution of MBW tracer gases in the lung, typically N_2 or sulfur-hexafluoride [1], a one-dimensional advection-diffusion transport equation for a scalar variable representing the normalized gas concentration $c = c(x,t) \in [0,1]$ was solved with a finite difference method (detailed description in S1 Appendix). Simulation of MBW involved the computation of gas concentration in all model airways over multiple breath periods. We considered only inert gases for which the dominant transport mechanisms are advection by the carrier gas (ambient air) and diffusion. The transport process was modelled along the centerlines of each pipe-like airway and along the axis of each lobule trumpet. The 1D approximation entails consideration of averaged (lumped) quantities within a cross-section, or in case of the trumpet model, within all airways represented by the cross-section of the trumpet at a given axial position. Advection-diffusion processes in a pipe-like geometry are subject to high radial velocity gradients, which can lead to radial concentration gradients such that the average effective diffusivity within a cross-section increases. In the human lung, these phenomena can take place at different scales: In the upper airway, where the *Reynolds* number Re_d (based on the airway diameter d) is about 10^4 , turbulent flow strongly enhances mixing. In smaller airways ($Re \ll 2000$), *Poiseuille* flow with a parabolic velocity profile can be assumed. Enhanced diffusion due to high velocity gradients was modelled based on the concept of *Taylor dispersion*, where the local diffusion coefficient is a function of the local *Peclet* number [37],

$$\hat{D} = D \left(1 + \frac{1}{192} Pe^2 \right) \quad (\text{Eq 9})$$

where \hat{D} and D are the effective and molecular diffusion coefficients, respectively, and $Pe = ud/D$ is the *Peclet* number defined with the local mean velocity u and the local airway diameter d . This is an approximate measure. Taylor dispersion is a concept, which strictly applies only to developed flow in straight tubes. Hence, the modification of the diffusion coefficient (Eq 9) does not account for local changes in diffusivity due to secondary flow phenomena occurring at bifurcations and curved airways, and likely underestimates the level of diffusion enhancement.

The advection-diffusion equation for gas transport was solved separately in each airway, applying interface conditions at the bifurcation nodes to couple the transport between different airways. We considered a transport equation of the following general form

$$\frac{\partial(Sc)}{\partial t} + \frac{\partial F}{\partial x} = 0 \quad (\text{Eq 10})$$

with the flux

$$F = S_{ad}uc - S\hat{D}\frac{\partial c}{\partial x} \quad (\text{Eq 11})$$

Here, we distinguished between the total cross-section S and the cross-section S_{ad} in which advection with the carrier gas velocity $u(x,t)$ takes place. This assumption is important, because

the airway geometry becomes increasingly complex (i.e. non-tubular, alveolar ducts and alveolar trees) for smaller airways, and tracer gas advection does not occur in the entire lumen [35, 38, 39]. At bifurcations, the concentration flux was conserved by enforcing $\Sigma F = 0$. In S1 Appendix, a model for the trumpet lobule is derived which allows a more specific form of Eq 10 to be stated for the gas transport within trumpet lobules with non-constant cross-section. In addition, the finite difference scheme for spatial derivatives and time-integration method used for the numerical solution of Eq 10 are explained.

Structural and mechanical modifications of the model

The proposed multi-scale model was designed to study the effects of functional and structural inhomogeneities of the peripheral airways on the N_2 washout procedure. To illustrate the capabilities of the model, several parameters of the trumpet lobules were systematically modified. The parameters were defined in a way that their effect on lung geometry and mechanics was physically meaningful and intuitively clear:

1. **Lobule Compliance:** Lung compliance, as a measure of distensibility of the lung, is not uniform within the healthy lung in vivo [40]. Moreover, it is found impaired in many lung diseases, e.g. increased in emphysema, and decreased in pulmonary fibrosis [41–43]. We tuned tissue compliance with the parameter ϕ , which is a non-dimensional factor that has a stiffening or softening effect on the elastic pressure. In the following the meaning of ϕ is explained in more detail: We considered a mean tidal volume $V_{lb}^{TV} = TV/N_{lb}$ per lobule, where TV is the tidal volume, and N_{lb} is the total number of trumpet lobules. We then defined the nominal lobule volume range $[V_{lb}^0, V_{lb}^0 + V_{lb}^{TV}]$. Furthermore, we considered a reference pressure amplitude $p_{lb}^{TV} = 1500$ Pa, which corresponds to a typical pressure range due to pleural pressure changes [44, 45]. For a given pressure load p_{lb}^{TV} , the elastic component of the lobule mechanics can be considered stiff (reduced compliance) if its maximum dynamic volume is smaller than V_{lb}^{TV} ($\phi < 1$), and soft (increased compliance, $\phi > 1$) otherwise. Fig 3A shows the elastic pressure curve (Eq 7) for a lobule for different values of ϕ , which effectively modify the curve via the shape parameter β as

$$\beta = \frac{p_{lb}^{TV}}{e^{\gamma V_{lb}^0} (e^{\phi \gamma V_{lb}^{TV}} - 1)} \tag{Eq 12}$$

The remaining shape parameter γ accounts for the non-linearity of the elastic pressure curve (Fig 3B). It was computed numerically for all the lobules (using a minimization approach) such that the elastic pressure curve intersects with the point $(\frac{1}{4} p_{lb}^{TV}, V_{lb}^0 + \frac{3}{4} \phi V_{lb}^{TV})$ in the volume-pressure plane. This point was chosen because the resulting curve qualitatively lies between a linear ($\gamma \rightarrow 0$) and an almost right-angled curve ($\gamma \rightarrow \infty$).

2. **Lobule Volume:** Volume differences between the respiratory units (i.e. the anatomical lobules) but also within each unit have been described in healthy lungs [6, 46] and are more pronounced in obstructive lung disease [47]. We used the modification parameter θ to modify the residual volume of a trumpet lobule as $V_{lb,mod}^0 = \theta V_{lb}^0$, such that $\theta < 1$ resulted in reduced residual trumpet lobule size.
3. **Lobule Resistance:** Differences in airway resistance [34] between separate airspaces are known to affect the washout process [48]. Such differences were introduced via a modification of the pressure loss (see Eq 7) in the trumpet lobule $p_{diss,mod} = \tau R_{lb} Q_{lb}$, where the modification parameter $\tau > 1$ directly modifies the lobular flow resistance R_{lb} . This can be used to model the obstruction of airways in a lobule, e.g. due to mucus plugging.

Comparison of model simulations with in vivo MBW measurements in healthy children

The aim of the comparison of model results with in vivo data was to demonstrate that the model can relate microscale modifications at the lobular level to clinically observed MBW metrics (MBW washout envelope and phase III slope analysis [1]).

We used data from healthy adolescents recruited for lung function studies in the Inselspital Children’s University Hospital, Bern, Switzerland. N₂MBW measurements (N = 4) were collected according to the recent ERS/ATS consensus guidelines [1] using the ultrasonic flowmeter (Exhalyzer D, Eco Medics AG, Duernten, Switzerland) and the software provided by the manufacturer (Spioware 3.1.6) as previously described [49]. During the test, the subject sat in an upright position wearing a nose-clip and was asked to breathe regularly through a snorkel-like mouthpiece connected to a bacterial filter and a dead space reducer.

Results & discussion

We use the above described model of the whole lung to study the effects of structural and mechanical modifications on MBW outcomes.

Baseline configuration vs. modified lobular compliance

A baseline simulation has been performed with the parameter settings listed in Table 1, but without any added asymmetries with respect to lobular compliance, lobular residual size, or lobular resistance. We compared this reference simulation with results for modified lobular compliance where two regions (each accounting for 25% of all trumpet lobules) were modified using $\phi = 0.5$ and $\phi = 1.5$, respectively. Fig 4 shows the results for both simulations (baseline and compliance modifications). The washout curve (Fig 4C and 4D) can be analyzed in different ways: Each expiration starts with zero N₂ concentration (phase I), which corresponds to the washout of the dead space (the pipes in the model), and then a rise in N₂ concentration, which is first very quick (phase II) and later slow (phase III) [50–52]. Phase III is the part of the concentration-expired volume that corresponds to 50% to 95% of the expired volume per breath [1] (Fig 5). The end-expiratory N₂ concentration diminishes from one breath to the next as the N₂ washout progresses. The ratio of two subsequent end-expiratory N₂ concentration values, i.e. the decay of the washout curve envelope, provides information about the gas washout efficiency of the whole lung, and indirectly about the ventilation of lung compartments. For the baseline model configuration, this exponential decay is uniform (linear envelope graph in a semi-logarithmic plot over time, in Fig 4D). In the model with regions of variable lobular compliance, the decay is non-uniform and delays towards the end. This can be

Table 1. Parameter settings of the baseline model.

Breathing profile	Tidal volume	TV	[l]	0.5
	Breath period	T _B	[s]	3.2
	Inlet flow profile			Sine function
	Length of simulation			50 breaths
Model settings	Functional residual capacity	FRC	[l]	3.0
	Limit diameter pipe airways	d _{lim}	[mm]	1.8
	Number of lobules		[-]	304
	Pipe generations			7–11*

* The number of pipe generations is not uniform for all model lobules, due to the asymmetrical branching scheme.

<https://doi.org/10.1371/journal.pcbi.1007079.t001>

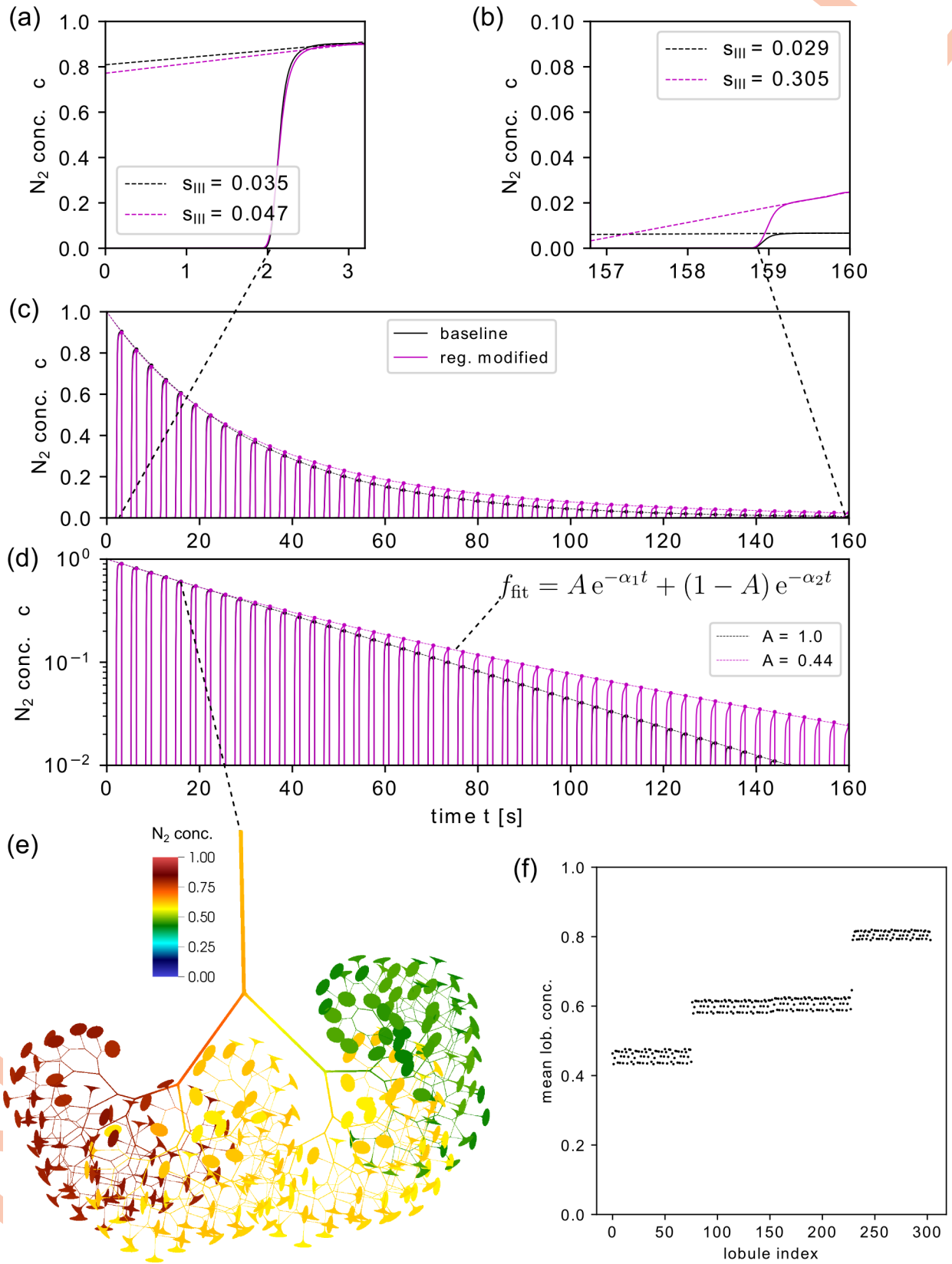


Fig 4. Overview of different simulation results for baseline model configuration and modified compliance. Results of the simulated N₂ gas washout for the baseline configuration (uniform and constant lung model parameter, black) and for altered trumpet lobule compliance using $\phi = 0.5, 1.5$ in two different regions, respectively, each accounting for 25% of all trumpet lobules (purple color). Normalized phase III slopes s_{III} are shown for the first (a) and last (b) breath. The washout profile (N₂ concentration at the entrance of the trachea) in (c) linear scale and (d) logarithmic scale for 50 simulated breaths with a uniform concentration decay for the baseline model, and a non-uniform decay for the regionally modified model (slow-fast washout profile). Panels (e) and (f) illustrate further the model with modified lobular compliance at the end of the fifth breath: (e) spatial N₂ concentration distribution where the trumpet lobules parametrized with lower and higher compliance are located on the left and right side of the airway tree, respectively. In (f) the mean lobular N₂ concentration is shown for each trumpet lobule.

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associated with a non-uniform ventilation in different regions of the lung. To quantify these washout properties, an exponential function of the following form was fitted to the envelope of end-expiratory values.

$$f(t) = Ae^{-\alpha_1 t} + (1 - A)e^{-\alpha_2 t} \tag{Eq 13}$$

Here, α_1 and α_2 (with $\alpha_1, \alpha_2 > 0$) are the decay rates of two superimposed processes, one fast and one slow [53], and $A \in [0, 1]$ represents the relative weight of the slow decay process ($\alpha_1 < \alpha_2$). The decay of the process is considered uniform if $A = 1$ and non-uniform if $A < 1$. Changing the compliance properties as described above resulted in $A = 0.44$, $\alpha_1 = 0.018$, and $\alpha_2 = 0.045$, compared to $A = 1.0$ and $\alpha_1 = 0.031$ for the baseline.

The spatial distribution within the lung model after the fifth breath (Fig 4E and 4F) provides further insight to the non-uniform (i.e. heterogeneous) washout process. The concentration remains high in lobules with decreased compliance ($\phi = 0.5$) and is reduced more rapidly in those with increased compliance ($\phi = 1.5$). Later in the washout process, the relative concentration difference between the two regions is therefore higher than in the baseline configuration and the washout slows down due to the increasingly dominant contributions of the slowly washed-out units.

The concentration profile can be further analyzed and interpreted on a per-breath basis. The slope of the concentration-volume curve during phase III (*slope III*) indicates whether the gas mixtures from different lung regions have different gas concentrations [35, 50] (Fig 5).

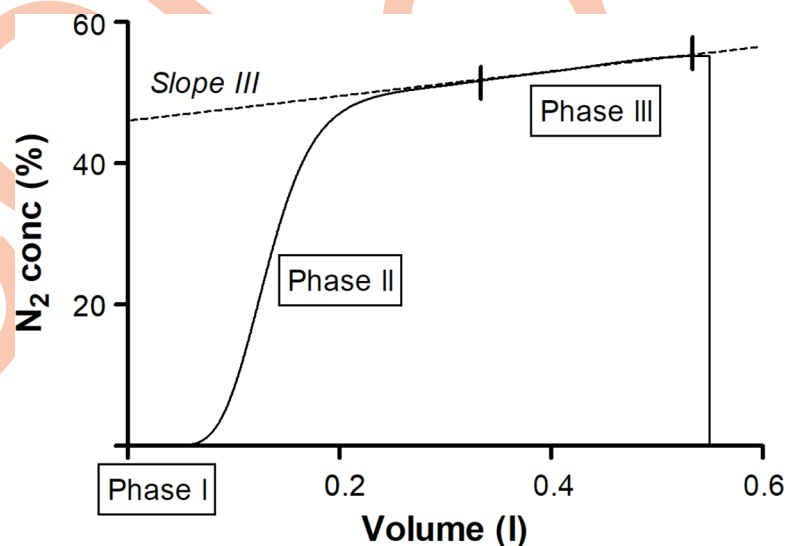


Fig 5. Schematic presentation of the N₂ concentration-volume curve of a single breath. Expired N₂ concentration is expressed as % of the initial N₂ concentration. Phase III is defined between 50% and 95% of the expired volume. Slope III: the slope of the concentration-volume curve during phase III.

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Therefore, *slope III* has been linked to inhomogeneous gas transport dynamics, and parameters derived from this analysis are of increasing clinical relevance [54–56]. To quantify the phase III profile in our study, the parameter s_{III} is defined, which is the slope of a linear function fitted to the N_2 concentration values corresponding to the phase III, normalized with the mean N_2 concentration during phase III (Fig 5):

$$S_{III} = \frac{\text{Slope III}}{\text{mean } N_2 \text{ (phase III)}} \quad (\text{Eq 14})$$

Comparing the baseline washout with the washout of modified compliance, s_{III} was higher (steeper slope) in the modified lung already for the first breath (Fig 4A), and this difference in s_{III} increased further until the last breath (Fig 4B).

Outcome parameters for comparison

For the comparison of other types of lung model modifications that represent a specific kind of structural and functional lung asymmetry, we used the weight parameter A and the decay rate α_1 (and α_2) from the fitting function (Eq 9) approximating the washout envelope, as well as the clinically important parameter s_{III} (Eq 14).

Other types of regional lobular modifications

In separate simulations, the residual volume of trumpet lobules was altered using $\theta = 0.5$ for a subset of lobules accounting for 25% of all trumpet lobules. Note that the model is designed such that it scales the residual volume of the remaining 75% of lobules automatically to preserve the prescribed FRC of the lung. In another simulation, the resistance was increased ($\tau = 8$) in 25% of the lobules, which corresponds to a local reduction of airway diameters by approximately 40%.

Fig 6 shows the simulation results for these three modified cases together with the baseline configuration. Note that the baseline results (black line in Fig 6) are barely visible, because some of the simulations in the model with increased resistance yielded very similar results. The compliance modifications showed the strongest effect and yielded a non-uniform, delayed washout (Fig 6C and 6D). While the resistance modification did not lead to any notable difference from the baseline result ($A = 1.0$), heterogeneous residual lobular volume also yielded a non-uniform washout ($A = 0.84$) similar but weaker than the model with modified compliance. Interestingly, the washout was delayed in later breaths in the case of modified residual lobular volume, and of compliance modifications. The modified residual volume caused in addition a faster decay for early breaths (Fig 6C before $t = 60$ s). This was also reflected in the slow ($\alpha_1 = 0.028$) and fast ($\alpha_2 = 0.074$) decay rates, e.g. when compared to the decay rate for the baseline configuration ($\alpha_1 = 0.031$). A reason for this could be that in smaller and shorter lobules, the flow of pure oxygen replaces a larger fraction of N_2 and thus the washout per breath is more efficient (faster). Although the residual lobule size is different, a similar flow rate of pure oxygen is preserved in all lobules as long as the compliance properties and the pleural pressure distribution remain uniform.

The comparison of normalized phase III slopes s_{III} also clearly discriminated the compliance modifications from the other cases. In a per breath analysis, a flat plateau-like profile resulted for the baseline, as well as for the cases with altered lobule size and resistance. For regionally altered lobule compliance, however, the gas concentration increased nearly linearly in phase III. This difference is already visible from the first breath, but becomes more prominent as the washout progresses (Fig 6A and 6B). For lobular resistance and volume modifications, the phase III concentration profile remained a flat plateau during the whole washout. In

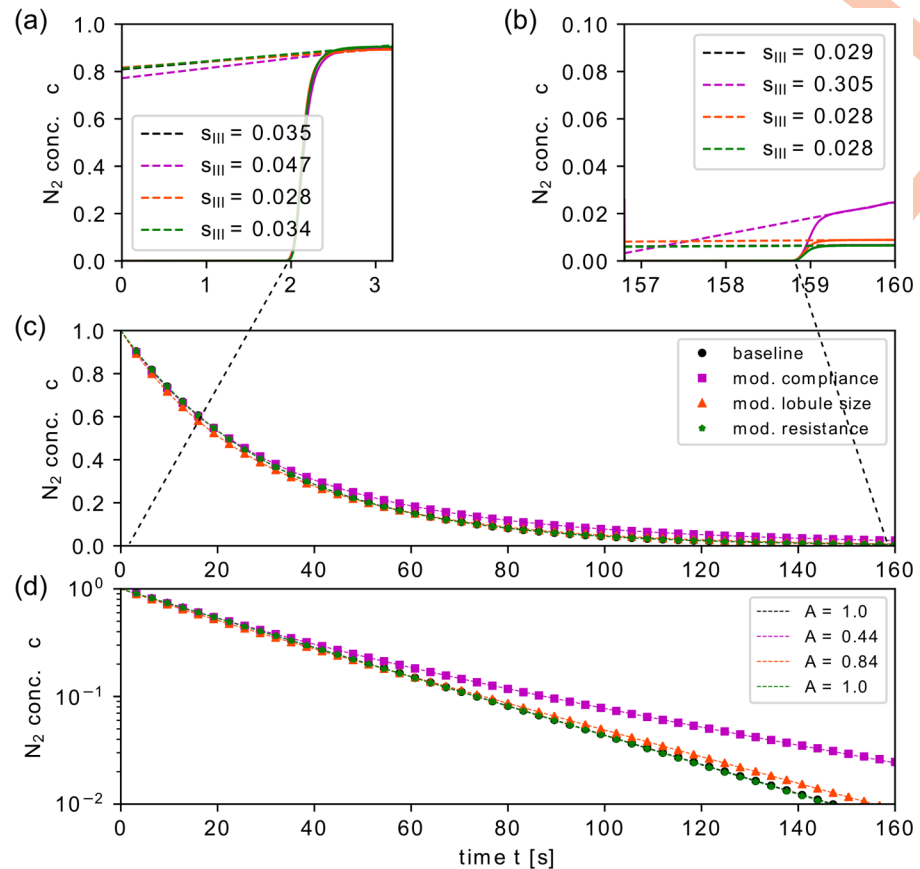


Fig 6. N_2 washout for baseline model configuration and modified trumpet lobule properties. Results of the simulated N_2 MBW for the baseline configuration (uniform and constant lung model parameters) and for three cases where different trumpet lobule parameters were modified regionally: 1) Lobular compliance was altered using $\phi = 0.5, 1.5$ in two regions (same as for results shown in Fig 4), 2) the lobular residual volume was decreased using $\theta = 0.5$, 3) the hydrodynamic resistance was amplified by a factor of 8 for 25% of all lobules. Normalized phase III slopes s_{III} are shown for the first (a) and last (b) breath. (c, d) The washout (only the envelope of the N_2 end-expiratory concentrations (symbols) and the corresponding fitting functions (dashed line, Eq 13) are shown) for 50 simulated breaths.

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the case of modified lobule compliance s_{III} increased from breath to breath. These different trends in s_{III} are illustrated in Fig 7. Increased slopes s_{III} indicate that concentration differences at airway bifurcations are increasing in phase III, such that contribution from high concentration units becomes more and more dominant. This was only the case for compliance modifications, where unequal rates of pure oxygen feed into different lobules. In case of modified volume differences, the spatial concentration distribution was not uniform, but did not change during phase III. In the next section, the relation between spatial concentration distribution in the lung and the phase III concentration profile is further discussed.

In summary, the compliance modifications had the strongest impact on gas washout in terms of non-uniform concentration decay and s_{III} values. A heterogeneity in residual lobule volume has a notable influence on the washout profile during early and later breaths, while the phase III concentration profile does not differ from the baseline. Compared to these two cases, the changes in hydrodynamic resistance have a negligible effect on gas washout profile. This is in accordance to a previous report [7].

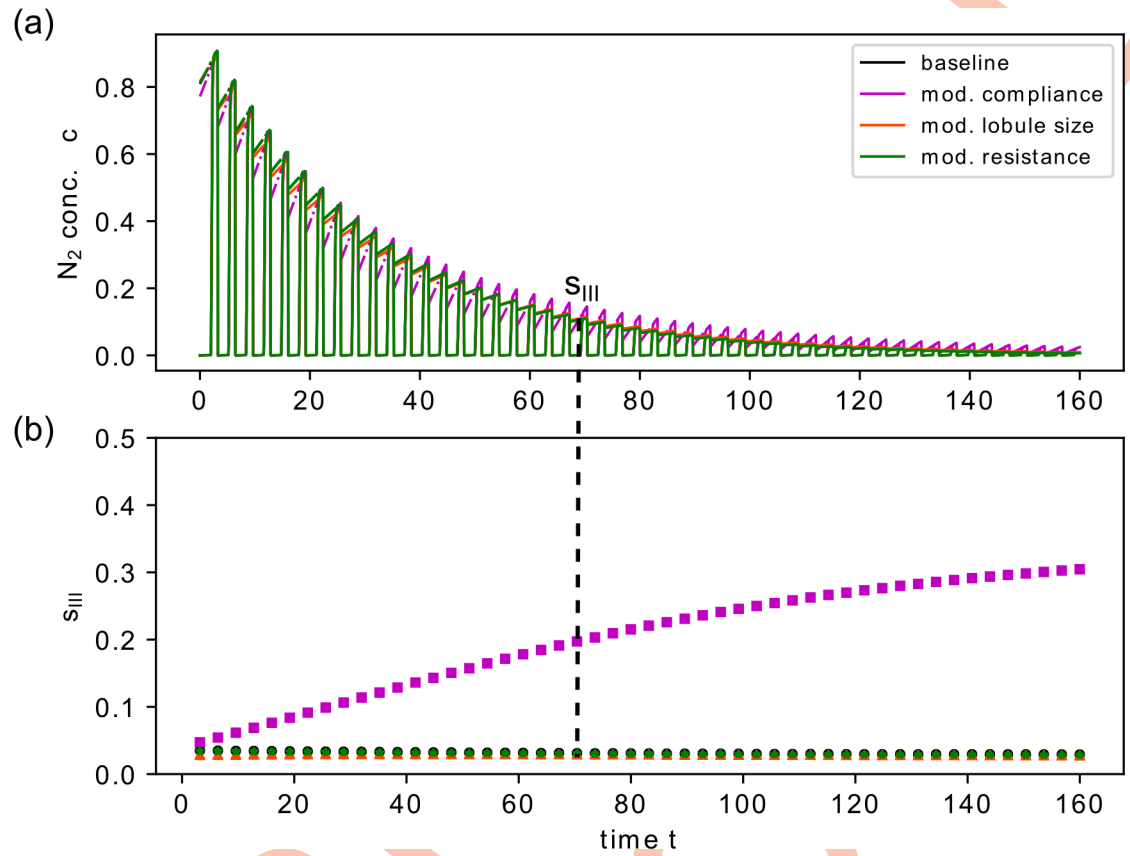


Fig 7. Phase III slopes s_{III} for baseline model configuration and modified trumpet lobe properties. Simulated N₂ washout profile for baseline and different types of regional modifications with phase III slopes s_{III} indicated for each breath (a). In (b) the values for the normalized phase III slopes for each breath are shown.

<https://doi.org/10.1371/journal.pcbi.1007079.g007>

Regional vs. local compliance modifications

To study the capabilities of the model in reflecting the effect of different spatial distributions of structural inhomogeneities, we look again at the example of compliance modifications. Instead of applying these modifications to a subset of neighboring lobules (regional distribution), we distributed the modifications over 50% of the lobules regularly distributed in the entire lung domain (local distribution). This distribution intends to mimic lung diseases such as cystic fibrosis that do not spread in a locally organized manner [57, 58]. To this end, we performed simulations where for every other lobule the compliance was alternately reduced ($\phi = 0.5$) or increased ($\phi = 1.5$).

Local and regional compliance modifications yielded similar results for the whole washout (Fig 8). Both caused a non-uniform decay with $A = 0.52$ (local) and $A = 0.44$ (regional). Minor differences were found in the slow and fast decay rates ($\alpha_1 = 0.020$ vs. 0.018 and $\alpha_2 = 0.044$ vs. 0.045 (local vs. regional)). However, the washout curve differed considerably with respect to phase III slope s_{III} (Fig 8A and 8B). Starting with only small differences in the first breath, s_{III} increased with every breath for regional modifications, while it decreased for local compliance modifications, reaching slightly negative values after approximately 35 simulated breaths (Fig 9D).

The spatial concentration distribution at the end of the fifth breath is depicted for both local and regional compliance modifications in Fig 9A and 9B. The lobular concentrations were clearly different for lobules with altered compliance properties. Trumpet lobules with

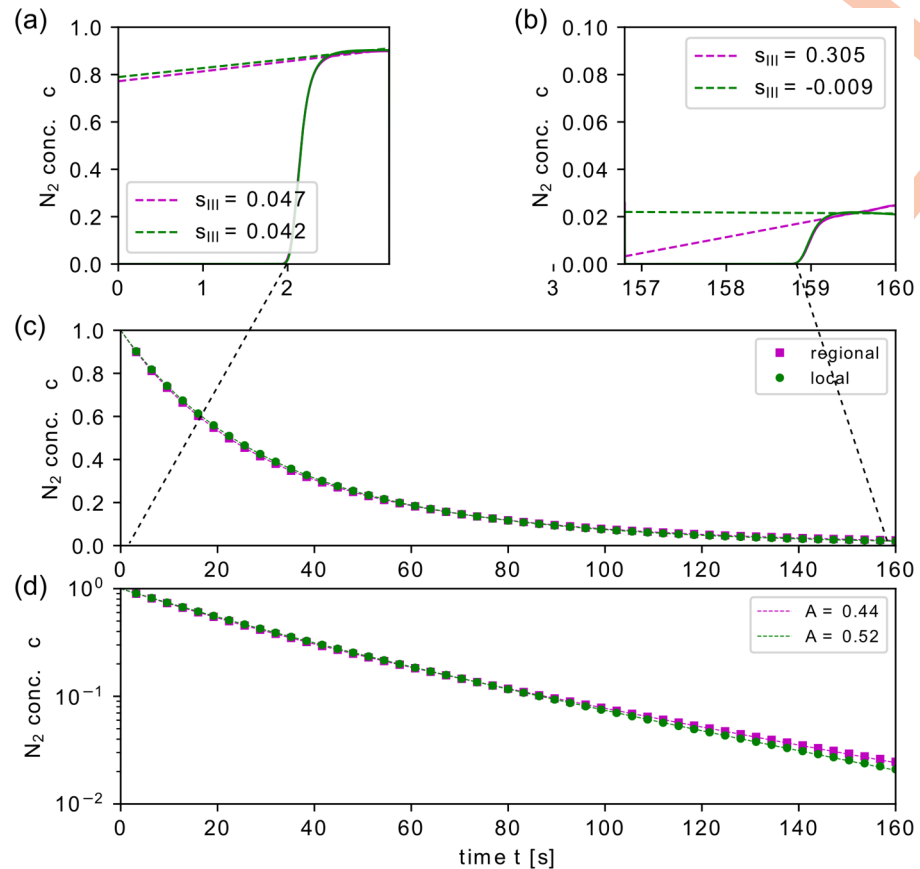


Fig 8. N_2 washout for regional and local distribution of compliance modifications. Results of the simulated N_2 gas washout for regional and local trumpet lobule compliance modifications. In both cases, the lobular compliance was altered using $\phi = 0.5$, and $\phi = 1.5$ each for 25% of all trumpet lobules. Normalized phase III slope s_{III} are shown for the first (a) and last (b) breath. The washout (c, d) for 50 simulated breaths, where only N_2 end-expiratory concentrations (symbols) and the corresponding fitting function (dashed line, Eq 13) are shown for better visibility.

<https://doi.org/10.1371/journal.pcbi.1007079.g008>

increased compliance properties showed higher (normalized) N_2 concentration and vice versa for reduced compliance. Furthermore, the comparison of the concentration distributions for regional and local compliance heterogeneity suggest the following mechanisms governing the phase III slopes: 1) The mixing of gas due to concentration gradients at airway bifurcations occurred at different levels: For regional modifications, the mixing took place at several generations of large airways, starting from the main bronchi to smaller conducting airways, where gas from different regions comes together, causing an increasingly non-uniform concentration distribution within the network. For local modifications, the mixing took place at already small airways and only over a few generations immediately above the lobules, which leads to a more uniform concentration distribution in the large airways. 2) Spatial concentration differences throughout the airway tree increased over time for regional modifications, while they remained approximately constant for local modifications (see S1 Video and S2 Video). This may be an explanation for the increasingly different phase III slopes (Fig 9C and 9D).

Model for the healthy lung

In order to mimic the small degree of inhomogeneous ventilation that exists physiologically in the healthy lungs [6], we used a combination of regional modifications of lobular size ($\theta = 0.5$

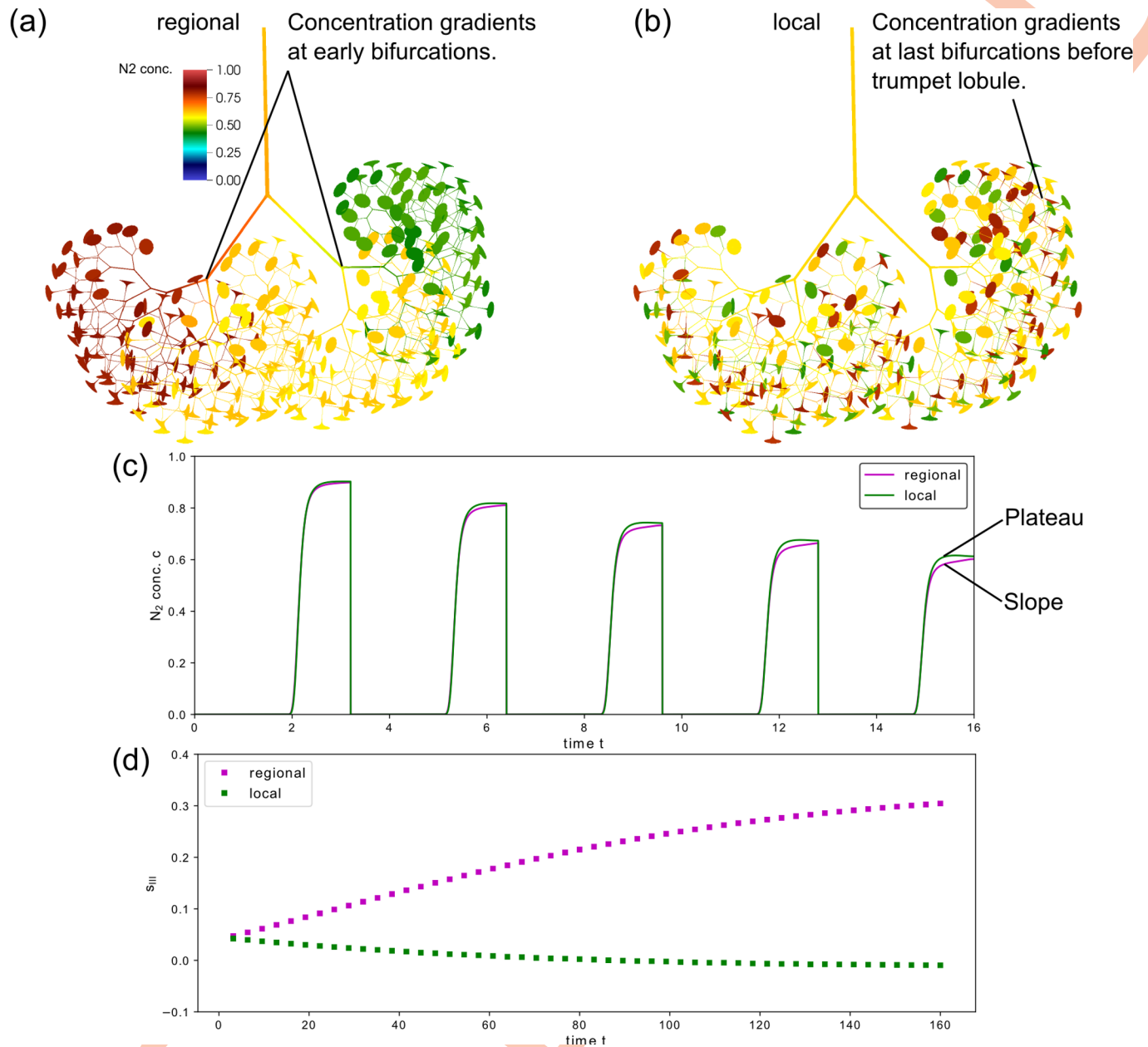


Fig 9. Spatial N_2 concentration distribution after the fifth breath for regional and local type compliance modifications. Spatial concentration distributions for regional (a) and local (b) type modifications at the end of the fifth breath. In trumpet lobules with lowered compliance ($\phi = 0.5$) the concentration remains high, whereas the lobules with increased compliance ($\phi = 1.5$) are washed out more efficiently. In panel (c), the corresponding washout profile is shown. The different profiles in the phase III (slope vs. plateau) for regional and local type modifications correlate with the pattern of the concentrations gradients at airway bifurcations. Panel (d) shows the temporal evolution (trend) of the normalized phase III slopes s_{III} for regional and local type modifications of the lobular compliance.

<https://doi.org/10.1371/journal.pcbi.1007079.g009>

in two regions each accounting for 12.5% of all trumpet lobules) and lobular compliance ($\phi = 0.5, \phi = 1.5$ in two regions respectively, each accounting for 12.5% of all trumpet lobules) to approximate normal structural and mechanical inhomogeneities of the lung. The modifications parameters were determined through an iterative process: First, the total set of lobules was split in sub-regions (each 12.5%) accounting for modified lobular compliance and size. From the previously conducted modification parameter study it was known that a region with

Table 2. Demographic characteristics and lung volumes for four MBW tests from healthy control adolescents.

MBW test	Subject	Age (years)	Weight (kg)	Height (cm)	sex	FRC (l)	VT (l)
#1	A	15.9	70.6	172.7	F	3.00	0.88
#2	B	17.9	50.0	169.5	M	2.85	0.67
#3	C	15.7	67.5	187.5	M	3.75	1.10
#4	C	15.7	67.5	187.5	M	3.52	0.63

M: male, F: female, FRC: functional residual capacity, VT: tidal volume. Of note, tests #3 and #4 were performed in sequence by the same subject (C).

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smaller lobule volume ($\theta < 1$) would account for faster washout during early breaths, while a region of less compliant lobule ($\phi < 1$) would cause the washout to be delayed (less efficient) towards later breaths. In a first step, the modification parameter θ was altered until the washout envelope of the measured and the simulated N_2 MBW curve were in agreement for the first 5–10 breaths (qualitatively, by visual inspection). Subsequently, the modification parameter ϕ was altered, which did not change the results from the first fitting step, until the washout envelopes also approximately matched for the last 30–50 breaths. In a MBW test simulation using this model, the lung clearance index (LCI) [1] was with 6.1 in the normal range reported in the literature [56]. [S3 Video](#) (supporting material) visualizes the evolution of the spatial concentration distribution over 50 breaths in such a model for a sinusoidal inlet flow profile. It shows the asymmetric concentration distribution due to regional differences in lobule size and lobule compliance, and how these differences evolve over time.

Comparison between simulations and real MBW measurements

Demographics about the healthy subjects and lung volumes for each test simulation are provided in [Table 2](#). For each simulation, we used the model for the healthy lung and applied the flow rate profile and the FRC as measured during the MBW test. The FRC was used to scale each lung model, using [Eq 1](#) with the reference value $FRC_W = 3.5$ l [26]. [Fig 10](#) and [Table 3](#) show measured concentration curves and MBW outcomes, respectively, for N_2 MBW measured in healthy subjects together with corresponding simulations.

Overall, the simulated washout envelopes for the four cases were in good agreement with the experimental data. On a breath-per-breath basis, differences were more prominent. For example, end-expiratory concentration values were moderately different for several breaths. A higher variability was found during phase III. Possible reasons for this could be: the complex structural and mechanical asymmetries in the healthy lung, which are not sufficiently modelled with the types of inhomogeneities used in this study (lobular compliance, residual size, and resistance); the non-uniform breathing pattern of the healthy subjects; and measurement errors. A detailed explanation or justification of these discrepancies was beyond the scope of this study.

Limitations of the model

The presented model has several limitations. For the sake of simplicity, several physiological and anatomical features were either idealized or neglected. The trumpet lobules include the last generations of the conducting airways, in order to keep the computational costs reasonable. We acknowledge that the acinar geometric properties are not entirely simulated in the lobule and that parameters like the homothety ratio are, in reality, not uniform throughout the airways contained in the lobule. It was beyond the scope of this study to investigate the effects of different cross-sectional development in the acinar region on the MBW washout curve.

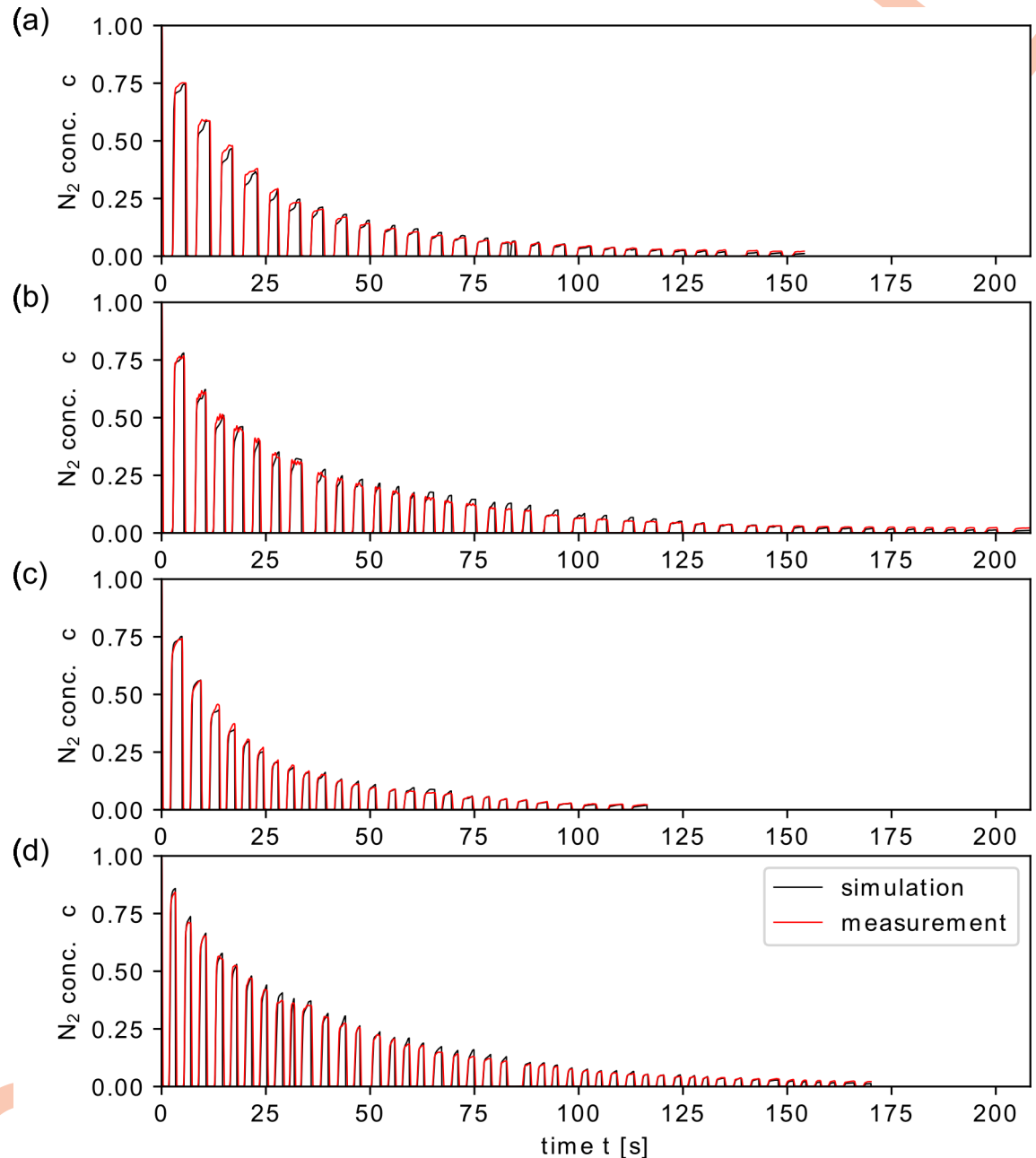


Fig 10. Measurements and simulation of N_2 MBW in healthy controls. Simulated N_2 washout compared to data from N_2 MBW tests from four healthy subjects (measurements (a)–(d)). For the simulations, the inlet flow profile and the FRC as measured during the N_2 washout test were used. To match the washout envelope in the measured data, both lobular residual volume and lobular compliance were partially modified in the lung model to mimic normal lung heterogeneity.

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Although the dimensions of the model (airway diameter and length per generation, FRC, lobular size distribution) were derived from anatomical data, the overall geometry of the model is rather generic and not anatomically based. Spatial asymmetries between the right and the left lung were not modelled. The FRC-dependent scaling of the modeled airway tree allows using it for MBW simulations in a large FRC range, however anatomical differences between children and adults were not introduced in the model.

Table 3. Multiple-breath washout (MBW) outcomes measured in reality and simulated in the lung model for four MBW tests from healthy control adolescents.

MBW test	LCI (TO) measured	LCI (TO) simulated	LCI washout breaths measured	LCI washout breaths simulated
#1	6.78	6.34	27	24
#2	6.37	6.22	31	30
#3	6.98	6.68	25	23
#4	7.12	6.75	43	40

LCI: lung clearance index, TO: turnovers. LCI washout breaths: number of washout breaths needed to reach the LCI

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Pressure losses due to turbulent flow in the big airways and flow changes at the bifurcations and/or the heavily curved airways are not included in the model, as these phenomena require 3D anatomical data. However, the results in this and in previous studies [7] demonstrate that flow resistance plays a smaller role in ventilation inhomogeneity compared to differences related to the loading and constitution of respiratory units. Although there is a spatial pleural pressure gradient in real lungs, the pleural pressure was assumed to be spatially uniform, because, to our best knowledge, there is no realistic estimate for this gradient. We further assumed that the initial mechanical stress distribution is uniform between the lobules although variability should be expected in real lungs. It will be interesting to investigate the effect of this variability and effects due to gravity on future studies.

Next, regarding the model for the healthy lungs, we acknowledge that various other combinations of θ and ϕ , as well as introduction of other modifying parameters may lead to a similar MBW profile and outcomes. However, our aim was neither to mimic breathing characteristics on the individual level [59], nor to relate MBW outcomes to particular local anatomical characteristics, but to provide an example that resembles healthy lungs, based on principles of lung physiology. Finally, no specific model for the upper airways, the mouth, and the interface equipment has been used in the simulations. The additional pathway for gas transport introduced by these parts was represented by an elongation of the straight pipe representing the trachea.

Conclusion

We presented a multi-scale model of the whole lung that simulates the gas transport and washout in conducting and acinar airways, including non-linear tissue mechanics. In order to mimic a physiological degree of ventilation inhomogeneity as described in healthy lungs, we introduced modifications in mechanical and geometrical properties on a lobular level. This study demonstrates that regional and local alterations of airway properties have different effects on the expiratory phase III in the MBW. Phase III slope profiles were notably more pronounced and sensitive to the degree of modifications for regional type modifications compared with local type modifications. Furthermore, the study revealed the different functional relations between the MBW concentration curves and airway compliance, volume and flow resistance. Increased heterogeneity of lobular compliance and residual volume correlated with a delayed washout, while heterogeneous flow resistance had a negligible impact. Finally, the simulation results are in accordance with real MBW data obtained from healthy subjects, on a qualitative level.

The model can be used to study MBW characteristics in health and disease. It offers the opportunity to understand the ventilation distribution in the healthy lung, and to investigate more profoundly MBW features that extract localized information, like the slope III analysis. By applying modifications in mechanical properties that exceed the physiological limits, the model can also mimic certain patterns of lung disease. Thus, it can be used to study the effect

of such diseases on MBW concentration curves. In addition, the model may also serve as a tool to visualize gas transport in the lung during a MBW test, which could support patient education.

Supporting information

S1 Appendix. 1. The trumpet lobule model. In this section, a geometry for the trumpet lobule cross-section is derived, which accounts for the cross-sectional growth due to exponentially increasing number of ever shorter airways, lumped into one unit (trumpet lobule). The resulting volume corresponds with the residual volume and the volume changes of a single lobule, which is prescribed by the ventilation LPM. Furthermore, a modified advection-diffusion equation is presented, which accounts for inert gas transport in the trumpet lobules with growing, time-varying cross-section. **2. Ventilation lumped parameter model.** In this section, the ventilation LPM is derived, which constitutes separate models for the non-compliant, pipe-like airways, and for the compliant (elastic) trumpet lobules. For both types of airway (unit) models the pressure-flow relation, respectively the pressure-volume relation are stated. The chosen methods for the numerical treatment of the LPM are also given in terms of an example based on a reduced size model. **3. Discretization of the transport equation.** Detailed information about the numerical treatment of the inert gas transport equation is provided. Spatial derivatives are approximated with second-order finite differences and for the temporal integration a Crank-Nicolson scheme is applied. A special focus is laid on the formulation used for an adequate treatment of the inert gas transport across airway bifurcations. Furthermore, the presented numerical scheme accounts for varying flow directions.

(PDF)

S1 Video. Evolution of spatial N₂-concentration distribution during first five breaths for regional compliance modifications. Same model modifications as in Fig 8.

(ZIP)

S2 Video. Evolution of spatial N₂-concentration distribution during first five breaths for local compliance modifications. Same model modifications as in Fig 8.

(ZIP)

S3 Video. Evolution of spatial N₂-concentration distribution during washout with 50 breaths for combined regional compliance modifications and regional size modifications. Same lung model modifications as for the model for the healthy lungs and the comparison with experimental data from healthy controls.

(ZIP)

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Visualization: David Hasler, Pinelopi Anagnostopoulou.

Writing – original draft: David Hasler, Pinelopi Anagnostopoulou.

Writing – review & editing: David Hasler, Pinelopi Anagnostopoulou, Sylvia Nyilas, Philipp Latzin, Johannes Schittny, Dominik Obrist.

References

1. Robinson PD, Latzin P, Verbanck S, Hall GL, Horsley A, Gappa M, et al. Consensus statement for inert gas washout measurement using multiple- and single- breath tests. *Eur Respir J*. 2013; 41(3):507–22. <https://doi.org/10.1183/09031936.00069712> PMID: 23397305
2. Subbarao P, Milla C, Aurora P, Davies JC, Davis SD, Hall GL, et al. Multiple-Breath Washout as a Lung Function Test in Cystic Fibrosis. A Cystic Fibrosis Foundation Workshop Report. *Ann Am Thorac Soc*. 2015; 12(6):932–9. <https://doi.org/10.1513/AnnalsATS.201501-021FR> PMID: 26075554
3. Boon M, Vermeulen FL, Gysemans W, Proesmans M, Jorissen M, De Boeck K. Lung structure-function correlation in patients with primary ciliary dyskinesia. *Thorax*. 2015; 70(4):339–45. <https://doi.org/10.1136/thoraxjnl-2014-206578> PMID: 25673230
4. Nyilas S, Schlegtendal A, Singer F, Goutaki M, Kuehni CE, Casaulta C, et al. Alternative inert gas washout outcomes in patients with primary ciliary dyskinesia. *Eur Respir J*. 2017; 49(1).
5. Marshall R. The physical properties of the lungs in relation to the subdivisions of lung volume. *Clin Sci*. 1957; 16(3):507–15. PMID: 13473164
6. Milic-Emili J. Ventilation Distribution. In: Hamid Q, Shannon J, Martin J, editors. *Physiologic Basis of Respiratory Disease*. Hamilton: BC Decker Inc; 2005. p. 133–42.
7. Swan AJ, Clark AR, Tawhai MH. A computational model of the topographic distribution of ventilation in healthy human lungs. *J Theor Biol*. 2012; 300:222–31. <https://doi.org/10.1016/j.jtbi.2012.01.042> PMID: 22326472
8. Bordas R, Lefevre C, Veckmans B, Pitt-Francis J, Fetita C, Brightling CE, et al. Development and Analysis of Patient-Based Complete Conducting Airways Models. *PLoS One*. 2015; 10(12):e0144105. <https://doi.org/10.1371/journal.pone.0144105> PMID: 26656288
9. Tawhai MH, Lin CL. Image-based modeling of lung structure and function. *J Magn Reson Imaging*. 2010; 32(6):1421–31. <https://doi.org/10.1002/jmri.22382> PMID: 21105146
10. Collier GJ, Kim M, Chung Y, Wild JM. 3D phase contrast MRI in models of human airways: Validation of computational fluid dynamics simulations of steady inspiratory flow. *J Magn Reson Imaging*. 2018; 48(5):1400–9. <https://doi.org/10.1002/jmri.26039> PMID: 29630757
11. Yin Y, Choi J, Hoffman EA, Tawhai MH, Lin CL. A multiscale MDCT image-based breathing lung model with time-varying regional ventilation. *J Comput Phys*. 2013; 244:168–92. <https://doi.org/10.1016/j.jcp.2012.12.007> PMID: 23794749
12. van Erbruggen C, Hirsch C, Paiva M. Anatomically based three-dimensional model of airways to simulate flow and particle transport using computational fluid dynamics. *J Appl Physiol* (1985). 2005; 98(3):970–80.
13. Ben-Tal A. Simplified models for gas exchange in the human lungs. *J Theor Biol*. 2006; 238(2):474–95. <https://doi.org/10.1016/j.jtbi.2005.06.005> PMID: 16038941
14. Foy BH, Kay D. A computational comparison of the multiple-breath washout and forced oscillation technique as markers of bronchoconstriction. *Respir Physiol Neurobiol*. 2017; 240:61–9. <https://doi.org/10.1016/j.resp.2017.02.016> PMID: 28300726
15. Henry FS, Llapur CJ, Tsuda A, Tepper RS. Numerical modelling and analysis of peripheral airway asymmetry and ventilation in the human adult lung. *J Biomech Eng*. 2012; 134(6):061001. <https://doi.org/10.1115/1.4006809> PMID: 22757498
16. Koshiyama K, Wada S. Mathematical model of a heterogeneous pulmonary acinus structure. *Comput Biol Med*. 2015; 62:25–32. <https://doi.org/10.1016/j.combiomed.2015.03.032> PMID: 25912985

17. Kumar H, Vasilescu DM, Yin Y, Hoffman EA, Tawhai MH, Lin CL. Multiscale imaging and registration-driven model for pulmonary acinar mechanics in the mouse. *J Appl Physiol* (1985). 2013; 114(8):971–8.
18. Fishler R, Hofemeier P, Etzion Y, Dubowski Y, Sznitman J. Particle dynamics and deposition in true-scale pulmonary acinar models. *Sci Rep*. 2015; 5:14071. <https://doi.org/10.1038/srep14071> PMID: 26358580
19. Mitchell JH, Hoffman EA, Tawhai MH. Relating indices of inert gas washout to localised bronchoconstriction. *Respir Physiol Neurobiol*. 2012; 183(3):224–33. <https://doi.org/10.1016/j.resp.2012.06.031> PMID: 22771781
20. Foy BH, Kay D, Bordas R. Modelling responses of the inert-gas washout and MRI to bronchoconstriction. *Respir Physiol Neurobiol*. 2017; 235:8–17. <https://doi.org/10.1016/j.resp.2016.09.009> PMID: 27677404
21. Verbanck S, Kerckx Y, Schuermans D, Vincken W, Paiva M, Van Muylem A. Effect of airways constriction on exhaled nitric oxide. *J Appl Physiol* (1985). 2008; 104(4):925–30.
22. Verbanck S, Paiva M. Model simulations of gas mixing and ventilation distribution in the human lung. *J Appl Physiol* (1985). 1990; 69(6):2269–79.
23. Verbanck S, Paiva M. A simulation study of diffusion-convection interaction and its effect on multiple breath washout indices. *Respir Physiol Neurobiol*. 2018; 258:5–11. <https://doi.org/10.1016/j.resp.2018.09.010> PMID: 30268738
24. Mountain JE, Santer P, O'Neill DP, Smith NMJ, Ciaffoni L, Couper JH, et al. Potential for noninvasive assessment of lung inhomogeneity using highly precise, highly time-resolved measurements of gas exchange. *J Appl Physiol* (1985). 2018; 124(3):615–31.
25. Majumdar A, Alencar AM, Buldyrev SV, Hantos Z, Lutchen KR, Stanley HE, et al. Relating airway diameter distributions to regular branching asymmetry in the lung. *Phys Rev Lett*. 2005; 95(16):168101. <https://doi.org/10.1103/PhysRevLett.95.168101> PMID: 16241843
26. Weibel ER. Principles and Methods of Morphometry. *Morphometry of the Human Lung*. Berlin, Heidelberg: Springer Berlin Heidelberg; 1963. p. 9–39.
27. Barre SF, Haberthur D, Cremona TP, Stampanoni M, Schittny JC. The total number of acini remains constant throughout postnatal rat lung development. *Am J Physiol Lung Cell Mol Physiol*. 2016; 311(6):L1082–L9. <https://doi.org/10.1152/ajplung.00325.2016> PMID: 27760763
28. Barre SF, Haberthur D, Stampanoni M, Schittny JC. Efficient estimation of the total number of acini in adult rat lung. *Physiol Rep*. 2014; 2(7).
29. Henninger HB, Reese SP, Anderson AE, Weiss JA. Validation of computational models in biomechanics. *Proc Inst Mech Eng H*. 2010; 224(7):801–12. <https://doi.org/10.1243/09544119JEIM649> PMID: 20839648
30. Florens M, Sapoval B, Filoche M. An anatomical and functional model of the human tracheobronchial tree. *J Appl Physiol* (1985). 2011; 110(3):756–63.
31. Miller W. *The Lung*. 2. ed. Springfield, Illinois: Charles C. Thomas; 1947.
32. Womersley JR. Oscillatory flow in arteries: effect of radial variation in viscosity on rate of flow. *J Physiol*. 1955; 127(2):38–9P. PMID: 14354701
33. Kaczka D, Colletti A, Tawhai M, Simon B. Computational Analyses of Airway Flow and Lung Tissue Dynamics. In: Chandran K, Udaykumar H, Reinhardt J, editors. *Image-Based Computational Modeling of the Human Circulatory and Pulmonary Systems* New York: Springer 2011. p. 375–402.
34. Bates J. *Lung mechanics*. Cambridge, UK: Cambridge University Press 2009.
35. Paiva M, Verbanck S. Gas convection and diffusion. In: Hamid Q, Shannon J, Martin J, editors. *Physiologic Basis of Respiratory Disease*. Hamilton, Ontario: BC Decker Inc; 2005. p. 143–54.
36. Verbanck S, Paiva M. Gas mixing in the airways and airspaces. *Compr Physiol*. 2011; 1(2):809–34. <https://doi.org/10.1002/cphy.c100018> PMID: 23737204
37. Cussler E. *Fundamentals of Diffusion*. Diffusion: Mass Transfer in Fluid Systems Cambridge: Cambridge University Press; 2009. p. 11–2.
38. Lovric G, Mokso R, Arcadu F, Vogiatzis Oikonomidis I, Schittny JC, Roth-Kleiner M, et al. Tomographic in vivo microscopy for the study of lung physiology at the alveolar level. *Sci Rep*. 2017; 7(1):12545. <https://doi.org/10.1038/s41598-017-12886-3> PMID: 28970505
39. Weibel ER, Sapoval B, Filoche M. Design of peripheral airways for efficient gas exchange. *Respir Physiol Neurobiol*. 2005; 148(1–2):3–21. <https://doi.org/10.1016/j.resp.2005.03.005> PMID: 15921964
40. Fry DL, Hyatt RE. Pulmonary mechanics. A unified analysis of the relationship between pressure, volume and gasflow in the lungs of normal and diseased human subjects. *Am J Med*. 1960; 29:672–89. PMID: 13702319

41. Christie RV. The Elastic Properties of the Emphysematous Lung and Their Clinical Significance. *J Clin Invest.* 1934; 13(2):295–321. <https://doi.org/10.1172/JCI100588> PMID: 16694212
42. Papandrinopoulou D, Tzouda V, Tsoukalas G. Lung compliance and chronic obstructive pulmonary disease. *Pulm Med.* 2012; 2012:542769. <https://doi.org/10.1155/2012/542769> PMID: 23150821
43. Plantier L, Cazes A, Dinh-Xuan AT, Bancal C, Marchand-Adam S, Crestani B. Physiology of the lung in idiopathic pulmonary fibrosis. *Eur Respir Rev.* 2018; 27(147).
44. Chiumello D, Chidini G, Calderini E, Colombo A, Crimella F, Brioni M. Respiratory mechanics and lung stress/strain in children with acute respiratory distress syndrome. *Ann Intensive Care.* 2016; 6(1):11. <https://doi.org/10.1186/s13613-016-0113-0> PMID: 26847436
45. Jonson B, Svantesson C. Elastic pressure-volume curves: what information do they convey? *Thorax.* 1999; 54(1):82–7. <https://doi.org/10.1136/thx.54.1.82> PMID: 10343639
46. Boyden EA. The structure of the pulmonary acinus in a child of six years and eight months. *Am J Anat.* 1971; 132(3):275–99. <https://doi.org/10.1002/aja.1001320302> PMID: 5115520
47. Ferguson GT. Why does the lung hyperinflate? *Proc Am Thorac Soc.* 2006; 3(2):176–9. <https://doi.org/10.1513/pats.200508-094DO> PMID: 16565428
48. Macklem P. Act of Breathing: Dynamics. In: Hamid Q, Shannon J, Martin J, editors. *Physiologic Basis of Respiratory Disease.* Hamilton: BC Decker Inc; 2015. p. 35–48.
49. Singer F, Houltz B, Latzin P, Robinson P, Gustafsson P. A realistic validation study of a new nitrogen multiple-breath washout system. *PLoS One.* 2012; 7(4):e36083. <https://doi.org/10.1371/journal.pone.0036083> PMID: 22558338
50. Verbanck S, Schuermans D, Van Muylem A, Paiva M, Noppen M, Vincken W. Ventilation distribution during histamine provocation. *J Appl Physiol* (1985). 1997; 83(6):1907–16.
51. Bashoff MA, Ingram RH Jr., Schilder DP. Effect of expiratory flow rate on the nitrogen concentration vs. volume relationship. *J Appl Physiol.* 1967; 23(6):895–901. <https://doi.org/10.1152/jappl.1967.23.6.895> PMID: 6065061
52. Horsfield K, Cumming G. Functional consequences of airway morphology. *J Appl Physiol.* 1968; 24(3):384–90. <https://doi.org/10.1152/jappl.1968.24.3.384> PMID: 5640725
53. Gustafsson PM, Robinson PD, Gilljam M, Lindblad A, Houltz BK. Slow and fast lung compartments in cystic fibrosis measured by nitrogen multiple-breath washout. *J Appl Physiol* (1985). 2014; 117(7):720–9.
54. Green K, Buchvald FF, Marthin JK, Hanel B, Gustafsson PM, Nielsen KG. Ventilation inhomogeneity in children with primary ciliary dyskinesia. *Thorax.* 2012; 67(1):49–53. <https://doi.org/10.1136/thoraxjnl-2011-200726> PMID: 21953064
55. Nyilas S, Bigler A, Yammine S, Kieninger E, Rochat I, Ramsey K, et al. Alternate gas washout indices: Assessment of ventilation inhomogeneity in mild to moderate pediatric cystic fibrosis lung disease. *Pediatr Pulmonol.* 2018; 53(11):1485–91. <https://doi.org/10.1002/ppul.24149> PMID: 30168294
56. Verbanck S, Paiva M, Schuermans D, Hanon S, Vincken W, Van Muylem A. Relationships between the lung clearance index and conductive and acinar ventilation heterogeneity. *J Appl Physiol* (1985). 2012; 112(5):782–90.
57. Nyilas S, Bauman G, Sommer G, Stranzinger E, Pusterla O, Frey U, et al. Novel magnetic resonance technique for functional imaging of cystic fibrosis lung disease. *Eur Respir J.* 2017;50(6).
58. Pennati F, Roach DJ, Clancy JP, Brody AS, Fleck RJ, Aliverti A, et al. Assessment of pulmonary structure-function relationships in young children and adolescents with cystic fibrosis by multivolume proton-MRI and CT. *J Magn Reson Imaging.* 2018; 48(2):531–42. <https://doi.org/10.1002/jmri.25978> PMID: 29457316
59. Dejours P, Bechtel-Labrousse Y, Monzein P, Raynaud J. [Study of the diversity of ventilatory rates in man]. *J Physiol (Paris).* 1961; 53:320–1.