Elisabet Åström Leif Uttman Lisbet Niklason Jerome Aboab Laurent Brochard **Björn Jonson**

Pattern of inspiratory gas delivery affects CO₂ elimination in health and after acute lung injury

Received: 30 November 2006 Accepted: 26 July 2007 Published online: 1 September 2007 © Springer-Verlag 2007

Electronic supplementary material The online version of this article (doi:10.1007/s00134-007-0840-7) contains supplementary material, which is available to authorized users.

E. Åström (🖂) · L. Uttman · L. Niklason · B. Jonson University Hospital, Department of Clinical Physiology, SE-221 85, Lund, Sweden e-mail: elisabet.astrom@med.lu.se Tel.: +46-46-173300 Fax: +46-46-151769 J. Aboab · L. Brochard

Hospital Henri Mondor, Medical Intensive Care Unit, 51, av. du Maréchal de Lattre de Tassigny, 94010 Créteil, France

Abstract Objective: To avoid ventilator induced lung injury, tidal volume should be low in acute lung injury (ALI). Reducing dead space may be useful, for example by using a pattern of inspiration that prolongs the time available for gas distribution and diffusion within the respiratory zone, the mean distribution time (MDT). A study was conducted to investigate how MDT affects CO₂ elimination in pigs at health and after ALI. Design and setting: Randomised crossover study in the animal laboratory of Lund University Biomedical Center. Subjects and intervention: Healthy pigs and pigs with ALI, caused by surfactant perturbation and lungdamaging ventilation were ventilated with a computer-controlled ventilator. With this device each breath could be tailored with respect to insufflation time and pause time $(T_{\rm I} \text{ and } T_{\rm P})$ as well as flow shape (square, increasing or decreasing flow). Measurements

and results: The single-breath test for CO₂ allowed analysis of the volume of expired CO₂ and the volume of CO₂ re-inspired from Y-piece and tubes. With a long MDT caused by long $T_{\rm I}$ or $T_{\rm P}$, the expired volume of CO2 increased markedly in accordance with the MDT concept in both healthy and ALI pigs. High initial inspiratory flow caused by a short $T_{\rm I}$ or decreasing flow increased the re-inspired volume of CO₂. Arterial CO₂ increased during a longer period of short MDT and decreased again when MDT was prolonged. Con*clusions:* CO_2 elimination can be enhanced by a pattern of ventilation that prolongs MDT. Positive effects of prolonged MDT caused by short $T_{\rm I}$ and decreasing flow were attenuated by high initial inspiratory flow.

Keywords Pulmonary gas exchange · Respiration, artificial · Capnography · Breath tests · Swine

Introduction

During mechanical ventilation, oxygenation can in most instances be maintained at very low alveolar ventilation by increasing the fraction of inspired oxygen. Exchange of CO_2 , however, depends upon alveolar ventilation. Enhanced CO₂ elimination without applying high airway pressure caused by high tidal volume (V_T) is an issue in some clinical situations. At acutely increased intracranial pressure a low P_aCO_2 at low airway pressures may be elimination, and an inspiratory pause may reduce respira-

desired, at least in an initial stage. In acute obstructive lung disease enhanced CO₂ elimination at low minute ventilation is often desired. Limitation of pressure and $V_{\rm T}$ is a strategy for lung-protective ventilation in acute lung injury (ALI) and acute respiratory distress syndrome (ARDS) [1–6]. With respect to gas exchange, an optimal benefit from a particular $V_{\rm T}$ is an issue, whatever target values for arterial pH or P_aCO_2 are chosen.

An optimal pattern of $V_{\rm T}$ delivery may enhance CO₂

tory dead space (V_D) or P_aCO_2 [7–11]. Such positive effects have not always been observed, however [9, 12–14]. Diverging results may reflect methodological limitations.

Capnography in the format of the single-breath test for CO_2 (SBT-CO₂) allows accurate determination of CO_2 elimination and partitions of V_D [15]. Uttman et al. showed in healthy pigs that CO₂ elimination depends on time available for gas distribution and diffusion within the respiratory zone, mean distribution time (MDT) [16]. Variation of MDT was achieved by changing the duration of the postinspiratory pause (T_P) . CO₂ elimination varied in proportion to the logarithm of MDT. Aboab et al. recently reported similar findings in ARDS patients [17]. In the latter study the concept of MDT was modified by taking into account the time required during inspiration to bring the fresh gas interface down to the respiratory zone. Furthermore, the concept of distribution was widened to include CO_2 exchange with alveolar blood. Still, knowledge is limited about how different combinations of duration of inspiratory gas insufflation $(T_{\rm I})$ and $T_{\rm P}$ affect CO₂ elimination and, ultimately, $P_{a}CO_{2}$.

The objective of the present study in pigs – healthy and after ALI – was to test the hypothesis that MDT describes how variation of inspiratory gas delivery affects breath-by-breath CO_2 elimination, when MDT is changed for one breath at a time. The study also explores whether a permanent lowering of MDT would constantly increase P_aCO_2 and vice versa.

Methods

Material

The local Ethics Board for Animal Research approved the study. Twenty-two pigs of the Swedish native breed, weighing 17–24 kg, were fasted overnight with free access to water. The animals were pre-medicated with azaperone (7 mg \cdot kg⁻¹) and anaesthetised with ketamine (5 mg \cdot kg⁻¹). Anaesthesia was maintained by continuous infusion of fentanyl (60 µg \cdot kg⁻¹ \cdot h⁻¹) and midazolam (0.7 mg \cdot kg⁻¹ \cdot h⁻¹).

ALI/ARDS was induced in 14 pigs by combining surfactant perturbation by inhalation of dioctyl sodium sulphosuccinate with very large $V_{\rm T}$ ventilation [18]. Details can be found in the electronic supplementary material (ESM) and in [19].

Pancuronium $(0.5 \text{ mg} \cdot \text{kg}^{-1} \cdot \text{h}^{-1})$ was given only to healthy pigs. In ALI pigs, paralysis was avoided in order to allow judgement of anaesthesia depth during the longer experiments. With the anaesthesia practised no muscular movements were observed. Ventilation was maintained using a 7.0 mm ID tracheal tube connected to a ventilator (ServoVentilator 900C, Siemens–Elema, Solna, Sweden). A mainstream analyser (CO₂ Analyzer 930, Siemens-Elema, Solna, Sweden) measured partial pressure of CO₂ at airway opening ($P_{ao}CO_2$). The ventilator/computer system used for data recording has previously been described [20]. Signals from the ventilator and CO₂ analyser representing flow rate, airway pressure and $P_{ao}CO_2$ were sampled at the frequency of 100 Hz. The signals had a 50% response time of 12 ms and were synchronous within ± 8 ms. Compliance of the tracheal tube and ventilator tubing was measured in vitro. The system was tested for leakage.

For all pigs, the ventilator was at baseline set at volume control with square inspiratory flow, $T_{\rm I}$ 33% and $T_{\rm P}$ 10% of the respiratory cycle. Minute ventilation was adjusted to achieve $P_{\rm a}\rm{CO}_2$ 5–6 kPa.

Protocol

Part 1: Inspiratory gas delivery modified for one breath at a time

Eight healthy pigs were, after preparation, stabilised for a period of 60 min at baseline ventilation at a fraction of inspired oxygen (F_IO₂) of 0.21. To combat the high tendency towards lung collapse in pigs, a positive endexpiratory pressure (PEEP) of 8 cmH₂O was used [21]. These pigs were studied at respiratory rate (RR) 20 min⁻¹ and 40 min⁻¹, denoted Health_{RR20} and Health_{RR40}. The non-linear influence of MDT on CO₂ exchange was considered to merit a primary exploration of particularly short MDT values at high RR, which may be used to reduce V_T in ARDS [19].

Six pigs were studied 24 h after induction of ALI/ARDS. These pigs were part of another study comparing how different modes of ventilation affect lung function (see ESM). After stabilisation at PEEP 10 cm H₂O and F_IO₂ 1.0, the effect of inspiratory flow patterns on CO₂ elimination of single breaths was studied at RR 20 min⁻¹. This group of pigs was denoted ALI_{RR20}.

For all groups of pigs (Health_{RR20}, Health_{RR40} and ALI_{RR20}) the pattern of inspiratory gas delivery was modified for single breaths at a time, with respect to $T_{\rm I}$ (0.2-1.9 s), $T_{\rm P}$ (0.1-0.5 s) and inspiratory flow wave form (SHAPE), in different combinations. SHAPE was either square, increasing, or decreasing flow rate. The latter two shapes were linear ramps starting or ending at zero flow. $V_{\rm T}$, PEEP and expiratory time were constant for all breaths. A recording sequence was pre-programmed in the computer that momentarily controlled the ventilator. Every 3rd breath out of 12 breaths comprising a recording sequence was modified. The breath immediately preceding a modified breath was defined as a control breath. Five different recording sequences, each with four modified breaths, gave 20 combinations of changes in $T_{\rm I}$, $T_{\rm P}$ and SHAPE. In randomised order, the five recording sequences were repeated three times.

Part 2: Prolonged periods of constant pattern of gas delivery

In eight healthy pigs (the same animals as the healthy pigs of part 1) and eight other pigs studied 4 h after induction of ALI/ARDS (ALI_{Tp}) alternative patterns of inspiratory gas delivery were maintained for prolonged periods in order to study the effect on P_aCO_2 . The reason for using a different group of ALI pigs than those used in part 1 was logistical; the total study time would otherwise have been too long. T_P was maintained at 17% for 40 min, then changed to 3%, which setting was again applied during 40 min, and finally the initial 17% was set again and applied also during 40 min. Thereby, during the middle period, MDT was changed by a factor of 0.5, i.e. from 80 ms to 40 ms. A reciprocal change in expiratory time maintained RR unchanged. Also V_T was unchanged. P_aCO_2 was measured every 5 min.

Data analysis

Sampled data of flow, pressure and $P_{ao}CO_2$ were transferred to an Excel workbook (Microsoft, Redmond, WA, USA) and analysed according to Uttman et al. [22]. Tidal CO₂ elimination (V_TCO_2) represents the difference between expired volume of CO₂ (V_ECO_2) and that reinspired from the Y-piece and adjacent tubing (V_ICO_2) (Fig. 1). Variations in V_ICO_2 , V_ECO_2 and V_TCO_2 resulting from variation of inspiratory pattern were expressed



Fig. 1 An example of single-breath test for CO₂ from a healthy pig. Expired volume of CO₂ ($V_{\rm E}$ CO₂), is the area under the expiratory curve (*grey and hatched areas*). Volume of CO₂ re-inspired from the Y-piece and adjacent tubing ($V_{\rm I}$ CO₂), is shown by the *hatched area*. The *grey area* represents tidal elimination of CO₂ ($V_{\rm T}$ CO₂). When inspiratory gas insufflation was prolonged from 0.75 s to 1.26 s, $V_{\rm E}$ CO₂ increased (*yellow area*) as a consequence of the increase in mean distribution time from 0.51 s to 0.66 s. The decrease in $V_{\rm I}$ CO₂ (*blue area*) was due to the decrease in $F_{\rm early}$ from 0.37 l/s to 0.121/s. Accordingly, $V_{\rm T}$ CO₂ increased by the sum of the yellow and blue areas

in percentage of average $V_{\rm T} \rm CO_2$ from the four control breaths in the same recording sequence immediately preceding the modified breaths and denoted $\Delta V_{\rm I} \rm CO_2$,

 $\Delta V_{\rm E} {\rm CO}_2$ and $\Delta V_{\rm T} {\rm CO}_2$, respectively. Airway dead space ($V_{\rm Daw}$) was defined as the point of maximum slope of the SBT-CO₂. Over the alveolar plateau, $P_{\rm ao} {\rm CO}_2$ was described by the equation

$$P_{\rm ao}\rm CO_2 = b + m \cdot \ln V_E \tag{1}$$

where $V_{\rm E}$ is volume of expired gas. Alveolar PCO₂ ($P_{\rm A}{\rm CO}_2$) represents the midpoint of the plateau.

Variation in $V_{\rm E}{\rm CO}_2$ results from a shift of the ascending limb along the volume axis that reflects a change in $V_{\rm Daw}$ ($\Delta V_{\rm Daw}$) and a shift of the alveolar plateau along the PCO₂ axis ($\Delta P_{\rm A}{\rm CO}_2$).

MDT was calculated from flow samples during $T_{\rm I}$ and $T_{\rm P}$ and with respect to $V_{\rm Daw}$ according to Aboab et al. [17]. Flow rate at onset of inspiration (F_{early}) was at square inspiratory flow $V_{\rm T}/T_{\rm I}$, at decreasing flow twice that value, and zero for increasing flow. The sum of $T_{\rm I}$ and $T_{\rm P}$ (T_{I+P}) was calculated as it represents time for transfer of CO₂ from blood to alveolar gas.

Statistical methods

Data are presented as mean \pm standard deviation (SD), or as mean \pm standard error of the mean (SEM) when error of the mean is the issue. Regression analysis was used to study variations in volumes of CO₂ in relation to parameters describing inspiratory flow pattern. Student's paired two-tailed *t*-test was used to analyse changes in P_a CO₂ during prolonged periods of altered T_P .

Results

 $V_{\rm T}$ and arterial blood gases are shown in Table 1. The ALI_{RR20} group was non-homogeneous with respect to gas exchange as reported in ESM.

Part 1

At Health_{RR20}, Health_{RR40} and ALI_{RR20}, $V_{I}CO_{2}$ for control breaths was 16 ± 1 , 34 ± 5 and $12 \pm 1\%$ of $V_{E}CO_{2}$,

Table 1 Tidal volume and arterial blood gases at the three conditions at start of measurements. In health PEEP was 8 and in ALI PEEP was 10 cm H_2O . Mean \pm SD

	V _T (ml/kg)	$P_{a}CO_{2}$ (kPa)	$P_{\rm a}O_2/F_{\rm I}O_2$ (kPa)
Health _{RR20}	8.8 ± 0.8	5.5 ± 0.5	63 ± 10
Health _{RR40}	5.9 ± 0.5	5.6 ± 0.3	62 ± 6
ALI _{RR20}	11.0 ± 1.7	5.9 ± 0.9	50 ± 17

$\Delta V_{\rm I} {\rm CO}_2 = a + b \cdot F_{\rm early}$			$\Delta V_{\rm E} \rm CO_2 = c + d \cdot \ln \rm MDT$		$\Delta V_{\rm T} \rm CO_2 = e + f \cdot \ln \rm MDT + g \cdot F_{early}$					
	a	b	R^2	С	d	R^2	е	f	g	R^2
Health _{RR20}	-3.3	19	0.67	9.0	18	0.70	12	20	-8	0.75
Health _{RR40}	-13	54	0.50	40	29	0.59	56	36	-27	0.71
ALI _{RR20}	-2.8	14	0.77	6.5	12	0.74	10	13	-13	0.81

Table 2 The relationship $\Delta V_{\rm I} \rm CO_2 = a + b \cdot F_{early}$, $\Delta V_{\rm E} \rm CO_2 = c + d \cdot \ln MDT$, and $\Delta V_{\rm T} \rm CO_2 = e + f \cdot \ln MDT + g \cdot F_{early}$ at square flow for the three conditions

 R^2 is correlation coefficient squared

Fig. 2 Variations in expired volume of CO₂, expressed in percentage of average tidal elimination of CO₂ from control breaths ($\Delta V_E CO_2$). $\Delta V_E CO_2$ related to mean distribution time (MDT) for all breaths studied and corresponding regression lines (black lines). Healthy pigs ventilated at low (Health_{RR20}) and high (Health_{RR40}) respiratory rate and pigs after induction of ALI/ARDS (ALI_{RR20}). Breaths with similar total inspiratory time (T_{I+P}) are indicated in a specific colour. Distributions around regression lines were independent of T_{I+P}



respectively. $\Delta V_{I}CO_{2}$ showed a significant positive correlation to F_{early} that reflects T_{I} and SHAPE (Table 2, Fig. 1).

In breaths with square flow but varying $T_{\rm I}$ and $T_{\rm P}$, $\Delta V_{\rm E} {\rm CO}_2$ increased significantly at higher MDT at all conditions. A logarithmic relationship was slightly better than a linear relationship for all conditions, but significantly so only for Health_{RR20} (p < 0.001) (Table 2, Fig. 2). Regression coefficient *d*, expressing the influence of lnMDT on $\Delta V_{\rm E} {\rm CO}_2$, was significantly higher at Health_{RR40} than at Health_{RR20} (p < 0.001). At ALI_{RR20} *d* was significantly lower than at Health_{RR20} (p < 0.001).

At Health_{RR20} and ALI_{RR20}, 56–58% of the change in $g \Delta V_{\rm E} {\rm CO}_2$ was caused by $\Delta V_{\rm Daw}$ and the remaining 42–44% by $\Delta P_{\rm A} {\rm CO}_2$. At Health_{RR40} the contribution to $g \Delta V_{\rm E} {\rm CO}_2$ by $\Delta V_{\rm Daw}$ was 69% and by $\Delta P_{\rm A} {\rm CO}_2$ 31%.

Differences between measured $\Delta V_{\rm E} {\rm CO}_2$ and values calculated from the logarithmic equations in Table 2 were calculated. No significant correlation between $T_{\rm I+P}$ and these residuals was found for Health_{RR20}, Health_{RR40} or ALI_{RR20}, as can be appreciated from Fig. 2. Accordingly, variations in $T_{\rm I+P}$ had no significant effect upon $\Delta V_{\rm E} {\rm CO}_2$ apart from effects explained by MDT.

At Health_{RR40}, when MDT was varied by changing SHAPE, $\Delta V_{\rm E} {\rm CO}_2$ varied similarly in relation to MDT as when $T_{\rm I}$ or $T_{\rm P}$ were varied. However, at Health_{RR20}, $\Delta V_{\rm E} {\rm CO}_2$ varied significantly less when MDT was changed by varying SHAPE. At ALI_{RR20}, no significant effect on $\Delta V_{\rm E} {\rm CO}_2$ was observed when SHAPE was varied. For breaths with square inspiratory flow, $\Delta V_{\rm T} {\rm CO}_2$



Fig.3 Observed P_aCO_2 when switching from a long to a short postinspiratory pause and back again, resulting in a change in mean distribution time from 0.8 s to 0.4 s. For each pig the values are normalised to the mean value of the three measurements just before shortening the postinspiratory pause. Average \pm SEM from six healthy pigs. During each period, the last three observations were used for calculation of data in Table 3

Part 2

At Health_{RR20} P_aCO_2 increased at short MDT (Fig. 3). The average of the three last observations during each period of 40 min was considered to represent steady state. At Health_{RR20} and ALI_{Tp}, average P_aCO_2 increased significantly during the period of short MDT and decreased significantly when MDT was again prolonged (Table 3). The change in P_aCO_2 was on average 75% of ΔV_TCO_2 resulting from changing MDT, estimated from equations in Table 2.

Discussion

The previously described system based upon a computercontrolled ventilator was amended to allow changed pattern of a single inspiration at a time [20]. In part 1, $T_{\rm I}$, $T_{\rm P}$ and SHAPE were modified while $V_{\rm T}$, expiratory time and PEEP were unchanged. This allowed a comprehensive analysis of how different patterns of inspiratory flow affect CO₂ exchange by using SBT-CO₂. By comparing modified breaths with control breaths in the same recording sequence, influence from even minor deviations from a steady state was avoided. The technique for modification of single breaths allowed studies of 21 inspiratory flow patterns in a short time. Uttman et al. introduced the concept of MDT to explain how inspiratory flow pattern affects CO₂ exchange by its effect on distribution of inspired gas in the alveolar zone [16]. Aboab et al. stressed that MDT refers not only to time for gas distribution and diffusion within alveolar space, but to time for all phenomena associated with transfer of CO2 from circulating blood in the alveolar capillaries to the airways [17]. This may include movements caused by the heart and pulsating blood.

 V_1 CO₂ was larger than the volume of CO₂ in the Y-piece connecting ventilator tubing to the airway. This reflects mixing of gas in the inspiratory and expiratory

Table 3 $\Delta P_a CO_2$ is the change
in PaCO2 after indicated change
in MDT expressed as percentage
of value before changing MDT
(see Fig. 3). $\Delta V_{\rm T} \rm CO_2$ is change
in $V_{\rm T} \rm CO_2$ resulting from the
change in MDT as estimated
according to equations in
Table 2. $\Delta P_a CO_2$ was on
average 75% of $\Delta V_{\rm T} \rm CO_2$.
Mean + SEM

		MDT $0.8 \text{ s} \rightarrow 0.4 \text{ s}$	MDT $0.4 \text{ s} \rightarrow 0.8 \text{ s}$	
Health _{RR20}	$\Delta P_{a}CO_{2}, \%$	8.8±2.8	-7.3 ± 1.5	
	$\Delta V_{\rm T} {\rm CO}_2$, % (estimated)	-11.8 ± 0.3	11.7±0.3	
ALI _{Tp}	$\Delta P_{\rm a} { m CO}_2, \%$	5.8±2.1	-5.1 ± 0.8	
	$\Delta V_{\rm T} {\rm CO}_2$, % (estimated)	-6.6±0.2	6.7±0.2	

lines shown by Fletcher et al. [23]. During the first part of inspiration, while CO₂ is present in both inspiratory and expiratory lines, a high flow rate increases turbulence and possibly also Coanda and Bernoulli effects around the Y-piece. This may explain the correlation between F_{early} and V_1CO_2 . V_1CO_2 amounted to about 16% of V_ECO_2 at Health_{RR20} and to 34% at Health_{RR40}. Re-inspiration of CO₂ is considerable. Within a low V_T strategy, reduction of V_1CO_2 can be achieved by one-way valves in the Y-piece, as suggested by Fletcher et al., or by aspiration of dead space, as discussed by De Robertis et al. [24].

In accordance with the hypothesis based upon previous studies, $\Delta V_{\rm E} \rm CO_2$ increased in relation to lnMDT [16, 17]. A better fit of a logarithmic equation rather than a linear one agrees with concepts based upon physiology and morphology. As gas distribution in lung periphery and exchange with alveolar blood depends on diffusion, gas exchange would be negligible at zero MDT. A very long MDT would imply that the interface between resident alveolar gas and fresh inspired gas is by diffusion brought up to a level at which the summed surface area according to the model of Weibel is so small that further diffusion becomes negligible [25]. When MDT increases from zero to high values, one may accordingly expect a fast initial increase in $\Delta V_{\rm E} \rm CO_2$ that becomes ever slower with further MDT increase. Patterns with particularly short MDT may severely reduce gas exchange, as the results at Health_{RR40} shows. When increased RR is used in ALI/ARDS in order to limit $V_{\rm T}$, it may be particularly important to maintain an adequate MDT by prolonging $T_{\rm P}$ and shortening expiration time.

At prolonged MDT, increasing $\Delta V_{\rm E} \rm CO_2$ reflected both a decrease of V_{Daw} and a positive $\Delta P_{\text{A}} \text{CO}_2$. This is in line with the results of Aboab et al. [17], who reasoned that a higher level of the alveolar plateau might, at least partially, be explained by continuing delivery of CO₂ by alveolar perfusion during a prolonged pause. T_{I+P} represents the time for alveolar perfusion during inspiration. In the present study we found that variation of T_{I+P} by different combinations of $T_{\rm I}$ and $T_{\rm P}$ did not significantly affect $\Delta V_{\rm E} {\rm CO}_2$ above what was explained by MDT. This suggests that time for alveolar perfusion during inspiration is of low importance compared to time for distribution and diffusion within the alveolar zone as expressed by MDT. A possible explanation why T_{I+P} in itself did not affect $\Delta V_{\rm E} {\rm CO}_2$ is that CO₂ delivered by alveolar perfusion late during inspiration does not, to a detectable extent, reach the upper respiratory zone in time to be expelled by the ensuing expiration.

A defined change in MDT had a similar effect on $\Delta V_{\rm E} {\rm CO}_2$ regardless of whether the change was caused by varying $T_{\rm I}$ or $T_{\rm P}$ (Fig. 2). Mathematical analysis shows that for a given increase in $T_{\rm P}$ the effect on MDT is three times larger than a comparable increase in $T_{\rm I}$ (see ESM). Accordingly, it is much more efficient to prolong $T_{\rm P}$ than $T_{\rm I}$.

The finding that at Health_{RR20} and at ALI_{RR20} , $\Delta V_{\rm E} {\rm CO}_2$ varied less and even insignificantly when MDT was changed by varying SHAPE implies that variation of SHAPE has effects on gas exchange that do not relate only to MDT. From studies based on flow oscillation techniques we know that sudden flow transients at airway opening lead to oscillations at frequencies around 5 Hz throughout the respiratory system. A hypothetical explanation for a maintained $\Delta V_{\rm E} \rm CO_2$ at increasing flow in spite of a shorter MDT is the following: At increasing flow the sudden end-inspiratory flow cessation leads to enhanced diffusion by way of important oscillations in lung periphery. Correspondingly, at decreasing flow, absence of end-inspiratory oscillations may explain why CO_2 elimination was lower than expected on the basis of prolonged MDT at this SHAPE.

In part 2, when MDT was varied for periods of 40 min, $P_{a}CO_{2}$ was expected to approach a steady state while CO_{2} stores in the body became equilibrated [26, 27]. During the periods of changed T_P and thereby changed MDT, $P_{\rm a}CO_2$ changed in the direction expected. The effect of reduction of T_P and MDT was less marked in ALI_{Tp} than in Health_{RR20}, but not significantly so. We do not speculate about the reasons for the possible difference observed. The change in $P_{a}CO_{2}$ was, for both Health_{RR20} and ALI_{Tp}, 75% of the change in $V_{\rm T} \rm CO_2$ estimated by using the equations in Table 2. In man, the time constant for the change in $P_{\rm a}{\rm CO}_2$ was 35 min when ventilation was decreased [27]. Incomplete steady state partially explains why the change in P_aCO_2 was < 100% of predicted change in V_TCO_2 . The period of changed $T_{\rm P}$ was limited to 40 min in order to limit interference from unavoidable changes in metabolism and other physiological factors affecting CO₂ exchange during prolonged experiments. Apart from incomplete steady state there may be other reasons why $P_{a}CO_{2}$ did not change as much as estimated change in $V_{\rm T}$ CO₂. For example, increasing intrapulmonary PCO₂ may affect distribution of pulmonary ventilation and perfusion or may change bronchial tone and thereby modify CO₂ elimination.

In Health_{RR20} the regression coefficient d was somewhat higher than in the non-homogeneous ALI_{RR20} group, indicating that an increase in MDT results in a slightly higher increase in CO_2 elimination in healthy pigs than in pigs after ALI. The present findings are comparable to observations in ARDS patients [17]. It appears that the absence of collateral ventilation in pigs has limited importance with respect to effects of MDT [28]. This is in line with the concept that a longer MDT promotes gas exchange by allowing more time for diffusion of gases in the border zone between conductive airways and the respiratory zone of the lung. Selection of an inspiration pattern providing a longer MDT can be made only in patients who are not breathing spontaneously. The improvement in gas exchange caused by a pattern optimising MDT may lead to reduction of $V_{\rm T}$ in the range of not more than 5–8% at ordinary RR. The non-linear relationship between CO₂

exchange and MDT implies that MDT becomes more important at increased rates. One should see optimisation of MDT as one way of reducing $V_{\rm T}$ which, in combination with other means of reducing dead space, may be important. One should also consider that reduction of dead space by any means paves the way for using higher RR. Importantly, the clinical implication of this study remains unclear until further studies have been performed in patients of different categories.

This study confirms that changes in pattern of inspiratory gas delivery lead to important instant changes in CO_2 elimination and ensuing changes in P_aCO_2 . At square inspiratory flow, these changes relate to MDT in accordance with the hypothesis. At increasing and decreasing

inspiratory flow rate, factors other than MDT must be further analysed. Flow rate early in inspiration affects the volume of re-inspired CO₂. The effects of changing MDT on CO₂ exchange are considerable, particularly at an increased RR. The findings merit further studies in critical care. The trade-off between improved gas exchange related to a prolonged MDT and potential negative effects of higher inspiratory flow rates and/or shorter expiration times should be investigated.

Acknowledgements. This study was supported by the Swedish Research Council (02872) and the Swedish Heart–Lung Foundation. Martina Christensson and Mikael Janiec performed complementary analyses of importance for interpretation of the results.

References

- Hickling KG, Henderson SJ, Jackson R (1990) Low mortality associated with low volume pressure limited ventilation with permissive hypercapnia in severe adult respiratory distress syndrome. Intensive Care Med 16:372–377
- Hickling KG, Walsh J, Henderson S, Jackson R (1994) Low mortality rate in adult respiratory distress syndrome using low-volume, pressure-limited ventilation with permissive hypercapnia: a prospective study. Crit Care Med 22:1568–1578
- Artigas A, Bernard GR, Carlet J, Dreyfuss D, Gattinoni L, Hudson L, Lamy M, Marini JJ, Matthay MA, Pinsky MR, Spragg R, Suter PM (1998) The American–European Consensus Conference on ARDS. Part 2: Ventilatory, pharmacologic, supportive therapy, study design strategies, and issues related to recovery and remodeling. Acute respiratory distress syndrome. Am J Respir Crit Care Med 157:1332–1347
- ARDSnetwork (2000) Ventilation with lower tidal volumes as compared with traditional tidal volumes for acute lung injury and the acute respiratory distress syndrome. N Engl J Med 342:1301–1308
- Petrucci N, Iacovelli W (2004) Ventilation with lower tidal volumes versus traditional tidal volumes in adults for acute lung injury and acute respiratory distress syndrome. Cochrane Database Syst Rev CD003844
- Kallet RH, Jasmer RM, Pittet JF, Tang JF, Campbell AR, Dicker R, Hemphill C, Luce JM (2005) Clinical implementation of the ARDS network protocol is associated with reduced hospital mortality compared with historical controls. Crit Care Med 33:925–929

- Fuleihan SF, Wilson RS, Pontoppidan H (1976) Effect of mechanical ventilation with end-inspiratory pause on blood–gas exchange. Anesth Analg 55:122–130
- Dammann JF, McAslan TC, Maffeo CJ (1978) Optimal flow pattern for mechanical ventilation of the lungs. 2. The effect of a sine versus square wave flow pattern with and without an end-inspiratory pause on patients. Crit Care Med 6:293–310
- Lachmann B, Jonson B, Lindroth M, Robertson B (1982) Modes of artificial ventilation in severe respiratory distress syndrome. Lung function and morphology in rabbits after wash-out of alveolar surfactant. Crit Care Med 10:724–732
- Wolff G, Brunner J, Weibel W, Bowes C (1989) Alveolar efficiency for CO₂ elimination and series dead space volume: both are affected by the ventilatory pattern. Appl Cardiopulm Pathol 2:309–314
- Mercat A, Diehl JL, Michard F, Anguel N, Teboul JL, Labrousse J, Richard C (2001) Extending inspiratory time in acute respiratory distress syndrome. Crit Care Med 29:40–44
- 12. Johansson H, Löfström JB (1975) Effects on breathing mechanics and gas exchange of different inspiratory gas flow patterns during anaesthesia. Acta Anaesthesiol Scand 19:8–18
- Al-Saady N, Bennett ED (1985) Decelerating inspiratory flow waveform improves lung mechanics and gas exchange in patients on intermittent positive-pressure ventilation. Intensive Care Med 11:68–75

- Markström A, Hedlund A, Lichtwarck-Aschoff M, Nordgren A, Sjöstrand U (2000) Impact of different inspiratory flow patterns on arterial CO₂ tension. Ups J Med Sci 105:17–29
- Beydon L, Uttman L, Rawal R, Jonson B (2002) Effects of positive end-expiratory pressure on dead space and its partitions in acute lung injury. Intensive Care Med 28:1239–1245
- Uttman L, Jonson B (2003) A prolonged postinspiratory pause enhances CO₂ elimination by reducing airway dead space. Clin Physiol Funct Imaging 23:252–256
- Aboab J, Niklason L, Uttman L, Kouatchet A, Brochard L, Jonson B (2007) CO₂ elimination at varying inspiratory pause in acute lung injury. Clin Physiol Funct Imaging 27:2–6
- Taskar V, John J, Evander E, Robertson B, Jonson B (1997) Surfactant dysfunction makes lungs vulnerable to repetitive collapse and reexpansion. Am J Respir Crit Care Med 155:313–320
- Uttman L, Ögren H, Niklason L, Drefeldt B, Jonson B (2007) Computer simulation allows goal-oriented mechanical ventilation in acute respiratory distress syndrome. Crit Care 11:R36
- Svantesson C, Drefeldt B, Sigurdsson S, Larsson A, Brochard L, Jonson B (1999) A single computer-controlled mechanical insufflation allows determination of the pressure-volume relationship of the respiratory system. J Clin Monit Comput 15:9–16
- 21. De Robertis E, Liu JM, Blomquist S, Dahm PL, Thorne J, Jonson B (2001) Elastic properties of the lung and the chest wall in young and adult healthy pigs. Eur Respir J 17:703–711

- 22. Uttman L, Jonson B (2002) Computeraided ventilator resetting is feasible on the basis of a physiological profile. Acta Anaesthesiol Scand 46:289–296
- 23. Fletcher R, Werner O, Nordström L, Jonson B (1983) Sources of error and their correction in the measurement of carbon dioxide elimination using the Siemens–Elema CO₂ Analyzer. Br J Anaesth 55:177–185
- 24. De Robertis E, Servillo G, Jonson B, Tufano R (1999) Aspiration of dead space allows normocapnic ventilation at low tidal volumes in man. Intensive Care Med 25:674–679
- 25. Weibel ER (1963) Morphometry of the human lung. Springer, Berlin, pp 110–143
- Farhi LE, Rahn H (1955) Gas stores of the body and the unsteady state. J Appl Physiol 7:472–484
- 27. Taskar V, John J, Larsson A, Wetterberg T, Jonson B (1995) Dynamics of carbon dioxide elimination following ventilator resetting. Chest 108:196–202
- Woolcock AJ, Macklem PT (1971) Mechanical factors influencing collateral ventilation in human, dog, and pig lungs. J Appl Physiol 30:99–115