

# Detection of optimal PEEP for equal distribution of tidal volume by volumetric capnography and electrical impedance tomography during decreasing levels of PEEP in post cardiac-surgery patients

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## Abstract

**Background:** Homogeneous ventilation is important for prevention of ventilator-induced lung injury. Electrical impedance tomography (EIT) has been used to identify optimal PEEP by detection of homogenous ventilation in non-dependent and dependent lung regions. We aimed to compare the ability of volumetric capnography and EIT in detecting homogenous ventilation between these lung regions.

**Methods:** Fifteen mechanically-ventilated patients after cardiac surgery were studied. Ventilator settings were adjusted to volume-controlled mode with a fixed tidal volume ( $V_t$ ) of 6–8 ml kg<sup>-1</sup> predicted body weight. Different PEEP levels were applied (14 to 0 cm H<sub>2</sub>O, in steps of 2 cm H<sub>2</sub>O) and blood gases,  $V_{cap}$  and EIT were measured.

**Results:** Tidal impedance variation of the non-dependent region was highest at 6 cm H<sub>2</sub>O PEEP, and decreased significantly at 14 cm H<sub>2</sub>O PEEP indicating decrease in the fraction of  $V_t$  in this region. At 12 cm H<sub>2</sub>O PEEP, homogenous ventilation was seen between both lung regions. Bohr and Enghoff dead space calculations decreased from a PEEP of 10 cm H<sub>2</sub>O. Alveolar dead space divided by alveolar  $V_t$  decreased at PEEP levels  $\leq 6$  cm H<sub>2</sub>O. The normalized slope of phase III significantly changed at PEEP levels  $\leq 4$  cm H<sub>2</sub>O. Airway dead space was higher at higher PEEP levels and decreased at the lower PEEP levels.

**Conclusions:** In postoperative cardiac patients, calculated dead space agreed well with EIT to detect the optimal PEEP for an equal distribution of inspired volume, amongst non-dependent and dependent lung regions. Airway dead space reduces at decreasing PEEP levels.

**Key words:** capnography; mechanical ventilation; peep; ventilator induced lung injury

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**Editor's key points**

- This pilot study compared volumetric capnography and electrical impedance tomography (EIT) in detecting distribution of ventilation after cardiac surgery.
- EIT values approximated to calculated dead space and detected optimal PEEP to equalize gas distribution in dependent and non-dependent lung regions.
- At 12 cm H<sub>2</sub>O PEEP, ventilation was distributed homogeneously between dependent and non-dependent regions.
- However, excessive PEEP applied to healthy lungs caused airway distension rather than improving alveolar ventilation.

Ventilator-induced lung injury (VILI) is a well-recognized complication of mechanical ventilation, which occurs in both patients with acute respiratory distress syndrome (ARDS)<sup>1</sup> and healthy lungs.<sup>2–3</sup> The deleterious effects of mechanical ventilation are caused by large amounts of stress and strain acting on lung tissue, as a result of inhomogeneous ventilation of the lungs.<sup>4–6</sup> Although, protective lung strategies using low tidal volumes have been strongly recommended to prevent the development of VILI, the amount of PEEP that should be applied is still under debate.<sup>7,8</sup> The positive effect of PEEP depends on the recruitability of lung tissue, which varies between patients,<sup>9–10</sup> but also from day to day. Therefore, setting the PEEP level without a reliable tool to estimate the distribution of inspiratory tidal volume (Vt) at the bedside is a challenging activity.

Volumetric capnography (Vcap) is a non-invasive technique that describes the CO<sub>2</sub> exhalation during one breath, and can be used to calculate alveolar and airway dead space.<sup>11</sup> Because elimination of CO<sub>2</sub> depends on alveolar ventilation and pulmonary perfusion, V/Q mismatches in either hyperinflated or collapsed lung areas will affect the amount of exhaled CO<sub>2</sub> and thereby Vcap. Recently, a multi-centre observational study showed that an increased dead space fraction in patients with ARDS was associated with an increased mortality.<sup>12</sup> Several studies showed the capability of Vcap in detecting optimal PEEP level during mechanical ventilation in various lung pathological conditions.<sup>13–15</sup> Böhm and colleagues<sup>14</sup> showed in 11 morbidly obese patients undergoing bariatric surgery, Slope III (S<sub>III</sub>) has a high sensitivity and specificity to detect lung recruitment. They concluded that S<sub>III</sub> was useful for identifying appropriate levels of PEEP in those patients. In 20 obese patients undergoing laparoscopic bariatric surgery Tusman and colleagues<sup>13</sup> found that both Vcap and pulse-oximetry, as compared with respiratory compliance, are accurate parameters for the detection of alveolar collapse. Fengmei and colleagues<sup>15</sup> found that best PEEP based on dead space fraction corresponded well with best PEEP according to highest respiratory compliance, in 23 ARDS patients. Therefore, Vcap is a promising technique to detect alveolar collapse and recruitment at the bedside.

Electrical Impedance Tomography (EIT) is a non-invasive, radiation-free, real time imaging modality, which has proved to correlate well with Computed Tomography (CT), according to assessment of changes in gas volume and tidal volume.<sup>16–18</sup> Recently, we showed that the intratidal gas distribution calculated from EIT measurements, is able to define a patient specific PEEP level, at which the lungs are homogeneously ventilated among dependent and non-dependent lung regions. This parameter was evaluated in both experimental studies and in patients,<sup>19–21</sup> but also during pressure support ventilation (PSV), Neurally Assisted Ventilatory Assist (NAVA) and pressure control ventilation

(PCV).<sup>22–23</sup> In addition, EIT has been used to visualize hyperinflation in the non-dependent region at a certain PEEP level when pixel-compliance is decreased.<sup>24–25</sup>

Suter and colleagues<sup>26</sup> defined optimal PEEP as the balance of adequate PaO<sub>2</sub> levels, good compliance and elimination of CO<sub>2</sub>. However, we believe that stress and strain are important contributors to lung injury, as both factors increase with inhomogeneous ventilation.<sup>5–6</sup> Therefore, the main goal of this study was to compare the results of Vcap measurements with that of EIT in finding the optimal PEEP, with an equal distribution of the inspiratory volume among the dependent and the non-dependent regions.

**Methods****Study population**

In this pilot-study, 15 mechanically ventilated patients, who had undergone coronary-artery bypass grafting and/or cardiac-valve surgery, admitted to the cardiothoracic intensive care unit (ICU) were included. The local medical ethical committee approved the study protocol and written informed consent was obtained from each patient or their relatives. Data were collected between January and July 2014.

The inclusion criteria were age >18 yrs, written informed consent, haemodynamically stable. Exclusion criteria were: presence of a cardiac pacemaker, pneumothorax, thoracic deformations and severe airflow limitation (defined as forced expiratory volume in 1 s below 70% of forced vital capacity).

**Study protocol**

For Vcap measurements a mainstream CO<sub>2</sub> sensor was placed between the tracheal tube and ventilator tubing's, and was connected to a NICO-capnograph (Novamatrix, Wallinford, Connecticut, USA). Specific software (Analysis plus, Novamatrix, Wallinford, Connecticut, USA) was used to record all Vcap data. In order to perform electrical impedance tomography (EIT) measurements, a silicon belt with 16 electrodes was placed around the thoracic cage between the 5th and 6th intercostal space (Pulmovista 500, Dräger, Lübeck, Germany). Ventilator settings were set to volume-controlled mode (Evita-infinity, Dräger, Lübeck, Germany), with a fixed tidal volume of 6–8 ml kg<sup>-1</sup> predicted body weight and inspiration/expiratory (I/E) ratio of 1:2. We deliberately choose volume control in order to avoid a change in the min volume because of differences in lung compliance at different levels of PEEP. The initial setting of the respiratory rate and fraction of inspired oxygen was adjusted to maintain an end-tidal carbon dioxide tension and oxygen saturation within a range of 35–45 mm Hg and 97–100%, respectively. PEEP was set according to the attending physician. Vt, respiratory rate (RR), I/E ratio remained unchanged throughout the entire study period.

We are reluctant to apply high levels of PEEP (>15 cm H<sub>2</sub>O) in postoperative patients after cardiac surgery who obviously have no ALI or ARDS, as these patients are generally prone to haemodynamic instability. Therefore, the patients were not subjected to recruitment manoeuvres. After baseline measurements of Vcap and EIT, the PEEP level was increased to 14 cm H<sub>2</sub>O. This PEEP level was maintained for 15–20 min in order to reach a steady state situation, as assessed by a stable signal of the volume of exhaled carbon dioxide (VCO<sub>2</sub>). Thereafter, the PEEP was decreased from 14 to 0 cm H<sub>2</sub>O PEEP in steps of 2 cm H<sub>2</sub>O. Each PEEP level was applied for five to 10 min depending on haemodynamically stability. Blood gas analysis, Vcap and EIT were measured during each PEEP step.

## Data analysis

### Volumetric capnography; model fitting and parameters

Vcap, which is constructed from expired CO<sub>2</sub> concentration and expired volume, can be represented by a mathematical function, which is previously introduced by Tusman and colleagues.<sup>27 28</sup> This makes the analysis of the Vcap data less susceptible to noise. To obtain this mathematical function from Vcap data, a non-linear least square curve fitting algorithm according to Tusman and colleagues was used in a custom-made Matlab® (The MathWorks, Natick, MA, USA) program. From this model, parameters were derived for further analysis. First, the Vcap curve was divided into three phases (Phase I, II and III) (Fig. 1). Phase I indicates the CO<sub>2</sub> volume originated from the airways, whereas phase III is the CO<sub>2</sub> from the alveoli. Phase II therefore is the phase with CO<sub>2</sub> from both the airways and alveoli. The inflection point of Phase II (Point A), which is determined by the maximum of the first derivative of the fitted model, was defined as the mean airway-alveolar interface that separates the airway from the alveolar compartment. Airway dead space (VD<sub>aw</sub>) is the volume from the beginning of the expiration until point A. Slope II (S<sub>II</sub>) is the slope of the inflection point of Phase II. Slope III (S<sub>III</sub>) is the slope of phase III, which is calculated according to Tusman and colleagues.<sup>27 28</sup> Therefore, phase<sub>III</sub> was divided into three even sized segments. A line was fitted through the second segment of phase<sub>III</sub> by the least-square method. The slope of this fitted line was determined as S<sub>III</sub>. Thereafter slope III was divided by the fraction of end tidal CO<sub>2</sub> (FECO<sub>2</sub>) in order to

normalize the value and make it comparable between patients.<sup>28 29</sup> Normalized Slope III (SnIII) represents the homogeneity of ventilation and pulmonary perfusion, which is considered as a good indicator of the global V/Q matching. SnIII increases in lung conditions associated with a mismatch of ventilation and perfusion such as atelectasis or pulmonary artery embolism, whereas low values indicate homogeneous V/Q matching.

Physiological dead space was calculated according to the formula developed by Bohr, and the Enghoff modification of Bohr's formula<sup>30</sup> (formula 1+formula 2):

(Mean alveolar partial pressure of CO<sub>2</sub> (P<sub>A</sub>CO<sub>2</sub>); mixed expired partial pressure of CO<sub>2</sub> (P<sub>E</sub>CO<sub>2</sub> =))

$$VD_{Bohr}/VT = \frac{P_A CO_2 - P_E CO_2}{P_A CO_2} \quad (1)$$

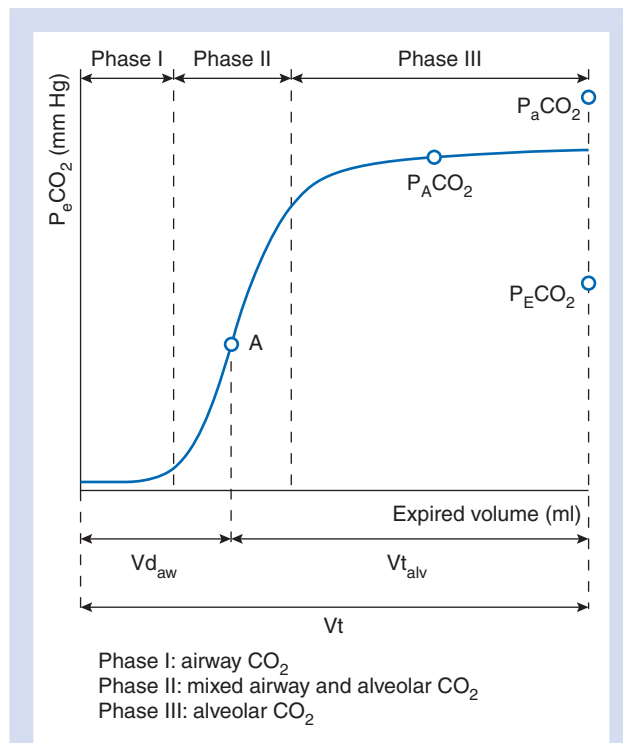
$$VD_{Enghoff}/VT = \frac{P_{aCO_2} - P_E CO_2}{P_{aCO_2}} \quad (2)$$

Alveolar dead space (VD<sub>alv</sub>) was computed by subtracting VD<sub>aw</sub> from VD<sub>Enghoff</sub>. We chose to subtract VD<sub>aw</sub> from VD<sub>Enghoff</sub> as the VD<sub>Enghoff</sub> contains all causes of V/Q inhomogeneity, whereas the Bohr formula is not supposed to contain shunts and low V/Q regions. The ratio of VD<sub>alv</sub> to alveolar tidal volume (VT<sub>alv</sub>) was obtained by dividing VD<sub>alv</sub> by VT<sub>alv</sub> (VT<sub>alv</sub>=VT-VD<sub>aw</sub>), in which VT<sub>alv</sub> represents alveolar tidal volume.

### EIT parameters

EIT data were recorded with a sample frequency of 20 Hz. Data was reconstructed and analysed using special software (EITdiag, Dräger, Lübeck, Germany). For each PEEP level, 10 to 20 consecutive breaths at the end of each PEEP step were selected for analysis, with the assumption that the lungs were best adapted to the ventilator settings at that time. EIT data contains signals induced by the respiratory system and the circulatory system. Therefore, to filter out signals related to the circulatory system, EIT data was filtered using a low pass-filter set to 40 beats min<sup>-1</sup>. The reconstructed image was represented as a ventilation distribution map, which was generated by tidal impedance changes caused, by inspiration and expiration. These impedance changes are presented as tidal impedance variation (TIV) and have been shown to correlate well with gas volume changes, as measured by CT-scans.<sup>16–18</sup> The surface area of the distribution map was divided into two equal regions of interest (ROIs), to know the non-dependent and dependent lung regions (Supplementary data, Fig. S1). The surface area of the EIT map was kept equal for all PEEP steps.

The intratidal gas distribution (ITV) was developed by Löwhagen, and colleagues<sup>21</sup> (formula 3). The ITV describes the amount of the impedance distributed to each region of interest within one inspiration. For this calculation Löwhagen and colleagues<sup>21</sup> divided the inspiratory part of the global impedance curve into eight equal volume sections. In other words each part of the inspiration is 12.5% of the entire inspiration. Thereafter, they transposed the time needed for each section of the inspiration to the regional curves. In this way they were able to calculate the contribution of four different regions of interest to the inspiration, during a single breath. According to our previous publications<sup>19 20 22 31</sup> we calculated the ITV for two instead of four regions of interest. Using the ITV calculation one is able to analyse the ventilation homogeneity during the inspiration. In order to reliably calculate the ITV, the filtered EIT signals were re-sampled with a frame rate of 40 Hz. This step is important in



**Fig 1** Schematic model of three phases of the expiration. The Vcap curve is divided in three phases. Phase I represents exhaled CO<sub>2</sub> from the airways, whereas phase III represents exhaled CO<sub>2</sub> from the alveoli. Phase II is a mixed phase with CO<sub>2</sub> from both the airways and alveoli. Point A is the inflection point of phase II, which is the theoretical point where the airway compartment is separated from the alveolar compartment. VD<sub>aw</sub>, airway dead space, VT<sub>alv</sub>, alveolar tidal volume, VT, tidal volume.

order to divide the inspiratory part of the global TIV curve reliably into eight iso-volume sections (ITV, *Intratidal Gas Distribution*; TIV, *Tidal Impedance Variation*; ROI, *Region of Interest*; t, *iso-volume part*).<sup>21</sup>

$$\text{Fractional regional ITV}_{1-8} = \frac{\text{ITV}_{1-8} \text{ TIV}_{\text{ROI}}}{\text{ITV}_{1-8} \text{ TIV}_{\text{Global}}} \quad (3)$$

In addition to the capnography and EIT parameters we also calculated the dynamic compliance. Dynamic compliance was calculated by dividing the expiratory tidal volume by Peak Inspiratory Pressure (PIP) minus PEEP.

### Statistical analysis

Statistical analyses were carried out using SPSS 21 (Chicago, IL, USA). Data are presented as mean (SD) unless otherwise specified. Normal distribution of the data was tested using the Kolmogorov-Smirnov test, whereas homoscedasticity was tested using the Brown-Forsythe test. Changes in Vcap and EIT parameters during the entire protocol were tested using mixed linear models. Differences between two sequential PEEP steps were tested using the paired Student's t-test for normal distributed data, and using the Wilcoxon rank sum test for not normal distributed data. For all statistical test  $P < 0.05$  was considered statistically significant.

### Results

The initial physiological data, measured shortly before 14 cm H<sub>2</sub>O was applied, is summarized in Table 1. During the different PEEP levels, blood gases and dynamic compliance remained stable, down to a PEEP level of 2 cm H<sub>2</sub>O and decreased significantly at 0 PEEP (ZEEP) (Table 2).

Figure 2 shows the TIV for the dependent and non-dependent lung region. The 14 cm H<sub>2</sub>O PEEP step was chosen as reference, and the TIV of both lung regions together was set to 100% at 14 cm H<sub>2</sub>O. Therefore, at lower PEEP levels the sum of TIV of

both regions can be more or less than 100%, as compared with PEEP 14 cm H<sub>2</sub>O. At 14 cm H<sub>2</sub>O of PEEP, most of the ventilation was distributed to the dependent lung region and from PEEP level of 10 cm H<sub>2</sub>O and lower, the non-dependent lung region became predominant ventilated (Fig. 2). TIV in the dependent lung region decreased during each PEEP step reduction from a PEEP of 12 cm H<sub>2</sub>O, indicating less ventilation at lower PEEP levels as a result of collapse (Fig. 2). TIV in the non-dependent region had the highest value at 6 cm H<sub>2</sub>O of PEEP (Fig. 2). In addition, ventilation was evenly distribution among dependent and non-dependent regions at 12 and 10 cmH<sub>2</sub>O PEEP (Fig. 3).

Table 2 shows changes in different Vcap parameters.  $\text{VD}_{\text{Bohr}}/\text{VT}$  and  $\text{VD}_{\text{Enghoff}}/\text{VT}$  significantly decreased from a PEEP of 10 cm H<sub>2</sub>O and less as compared with 14 cm H<sub>2</sub>O PEEP. The ratio of alveolar dead space to alveolar tidal volume significantly decreased from a PEEP level of 6 cm H<sub>2</sub>O and less as compared with 14 cm H<sub>2</sub>O PEEP. The normalized slope of phase III significantly changed at a PEEP levels of 4 cm H<sub>2</sub>O or less, whereas volume of phase III corrected for tidal volume ( $V_{\text{III}}/\text{VT}$ ) increased significantly, as compared with 14 cm H<sub>2</sub>O PEEP at  $\leq 8$  cm H<sub>2</sub>O PEEP, indicating more alveolar ventilation at lower PEEP levels. In addition,  $\text{VD}_{\text{aw}}$  decreased significantly from a PEEP level of 8 cm H<sub>2</sub>O or less. Shunt fraction calculated as the difference between arterial CO<sub>2</sub> minus end-tidal CO<sub>2</sub> ( $\text{Pa}_{\text{ET}}\text{CO}_2$ )<sup>32</sup> only significantly differed at ZEEP.

Figure 4A shows the effect of PEEP on the Vcap curve. When lower PEEP levels are applied the Vcap curve is shifted to the left, as a consequence of decreased  $\text{VD}_{\text{aw}}$ .  $\text{VD}_{\text{aw}}/\text{VT}$  significantly decreased at 10 cm H<sub>2</sub>O and lower as compared with 14 cm H<sub>2</sub>O PEEP (Table 2).

### Discussion

In this study we found that dead space calculations according to Bohr and Enghoff agreed well with the PEEP level, resulting in

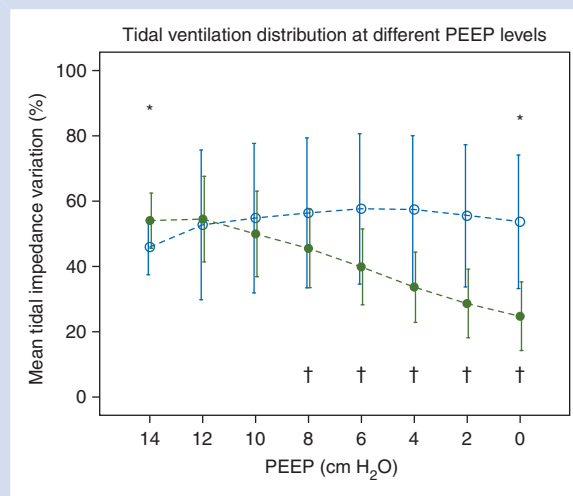
**Table 1** Data are presented as mean (SD) unless otherwise specified. Predicted body weight (PBW), Cardiopulmonary bypass (CPB), Coronary artery by-pass graft (CABG), Arterial partial pressure of O<sub>2</sub> ( $\text{Pa}_{\text{O}_2}$ ), Fraction of inspired oxygen ( $\text{Fi}_{\text{O}_2}$ ), Arterial partial pressure of CO<sub>2</sub> ( $\text{Pa}_{\text{CO}_2}$ )

Patient characteristics			
No. of patients			15
Age (yr)			70 (8)
Male/Female			13/2
Height (cm)			176 (13)
Weight (kg)			86 (18)
Predicted body weight (kg)			70 (13)
BMI			28 (4)
CPB time (min)			112 (35)
Type of surgery	CABG	Valve replacement	CABG+Valve replacement
	10	3	2
Haemodynamic data at ICU admission			
Mean arterial pressure (mm Hg)			70 (9)
Heart rate (BPM)			77 (15)
Ventilator settings and respiratory measurements at ICU admission			
Positive end-expiratory pressure (cm H <sub>2</sub> O)			8 (1)
Peak inspiratory pressure (cm H <sub>2</sub> O)			22 (2)
Expiratory tidal volume (ml)			494 (74)
Expiratory tidal volume/Predicted body weight (ml kg PBW <sup>-1</sup> )			7.1 (0.6)
Respiratory rate(Bpm)			17 (2)
$\text{Pa}_{\text{O}_2}/\text{Fi}_{\text{O}_2}$ (mm Hg)			350 (101)
$\text{Pa}_{\text{CO}_2}$ (mm Hg)			42.2 (4.5)



**Table 2** Data are presented as mean (SD). The first statistical change compared with 14 cm H<sub>2</sub>O PEEP is indicated by \*P<0.05 was considered significant. Tidal volume (VT), Airway dead space (VD<sub>aw</sub>), Alveolar dead space (VD<sub>alv</sub>), Alveolar dead space to alveolar tidal volume ratio (VD<sub>alv</sub>/VT<sub>alv</sub>), Normalized slope of phase III (S<sub>III</sub>), Amount of expired CO<sub>2</sub> within one breath (VTCO<sub>2,br</sub>), Volume of phase III to tidal volume ratio (VIII/VT), Arterial minus end-tidal partial pressure of CO<sub>2</sub> (Pa-ETCO<sub>2</sub>), Arterial partial pressure of O<sub>2</sub> (PaO<sub>2</sub>), Arterial partial pressure of CO<sub>2</sub> (PaCO<sub>2</sub>), Fraction of inspired oxygen (FiO<sub>2</sub>)

PEEP (cm H <sub>2</sub> O)	14	12	10	8	6	4	2	0
VT (ml)	472 (80)	476 (77)	474 (80)	473 (77)	472 (76)	471 (78)	474 (81)	473 (81)
VD <sub>aw</sub> (ml)	214 (40)	209 (42)	204 (39)	196 (40)*	188 (39)	182 (36)	174 (32)	167 (26)
VD <sub>alv</sub> (ml)	52 (37)	52 (34)	47 (31)	45 (33)	42 (33)*	35 (32)	39 (31)	35 (29)
VD <sub>Bohr</sub> /VT	0.48 (0.06)	0.47 (0.07)	0.45 (0.07)*	0.43 (0.07)	0.40 (0.07)	0.38 (0.07)	0.35 (0.07)	0.33 (0.07)
VD <sub>Enghoff</sub> /VT	0.57 (0.07)	0.55 (0.08)	0.53 (0.08)*	0.52 (0.08)	0.49 (0.08)	0.47 (0.09)	0.46 (0.08)	0.44 (0.08)
VD <sub>alv</sub> /VT <sub>alv</sub>	0.19 (0.11)	0.18 (0.1)	0.17 (0.09)	0.15 (0.09)	0.14 (0.1)*	0.12 (0.1)	0.13 (0.09)	0.11 (0.09)
S <sub>III</sub> (mm Hg ml <sup>-1</sup> )	0.95 (1.19)	0.73 (0.64)	0.62 (0.31)	0.62 (0.43)	0.58 (0.33)	0.48 (0.25)*	0.47 (0.26)	0.50 (0.21)
VTCO <sub>2,br</sub> (ml)	12.5 (3.3)	13.3 (3.7)*	13.9 (3.7)	14.3 (3.6)	14.7 (3.7)	15.2 (3.7)	15.7 (3.8)	16.0 (3.9)
VIII/VT	0.42 (0.14)	0.39 (0.13)	0.41 (0.14)	0.43 (0.13)*	0.46 (0.13)	0.46 (0.13)	0.48 (0.12)	0.51 (0.11)
Pa-ETCO <sub>2</sub> (mm Hg)	3.6 (2.8)	3.4 (2.5)	3.7 (2.6)	3.2 (2.5)	2.9 (2.7)	2.7 (2.4)	3.1 (2.4)	2.8 (2.5)*
PaCO <sub>2</sub> (mm Hg)	43 (5)	43 (5)	44 (5)	43 (5)	42 (5)	42 (5)	42 (5)	41 (5)*
PaO <sub>2</sub> (mm Hg)	138 (35)	140 (32)	145 (25)	145 (24)	145 (23)	139 (21)	132 (24)	127 (23)*
PaO <sub>2</sub> /FiO <sub>2</sub> (mm Hg)	346 (87)	349 (80)	359 (71)	371 (71)	372 (68)	357 (65)	338 (72)	324 (70)*
Dynamic compliance (ml cm H <sub>2</sub> O <sup>-1</sup> )	34 (6)	36 (8)	35 (6)	35 (7)	35 (7)	34 (7)	33 (7)	31 (7)*



**Fig 2** Tidal Impedance Variation at different PEEP levels. Data are shown as mean (SD). In the dependent lung region the ventilation distribution was decreased when PEEP was lowered, whereas the non-dependent region received more ventilation as compared with PEEP 14 cm H<sub>2</sub>O. \* Indicates a significant reduction in TIV of the non-dependent region according to 6 cm H<sub>2</sub>O. † Indicates a significant reduction in TIV of the dependent region as compared with 12 cm H<sub>2</sub>O. Dashed lines represents the interpolation lines; open circles=non-dependent region; solid circles=dependent region. P<0.05 was considered significant.

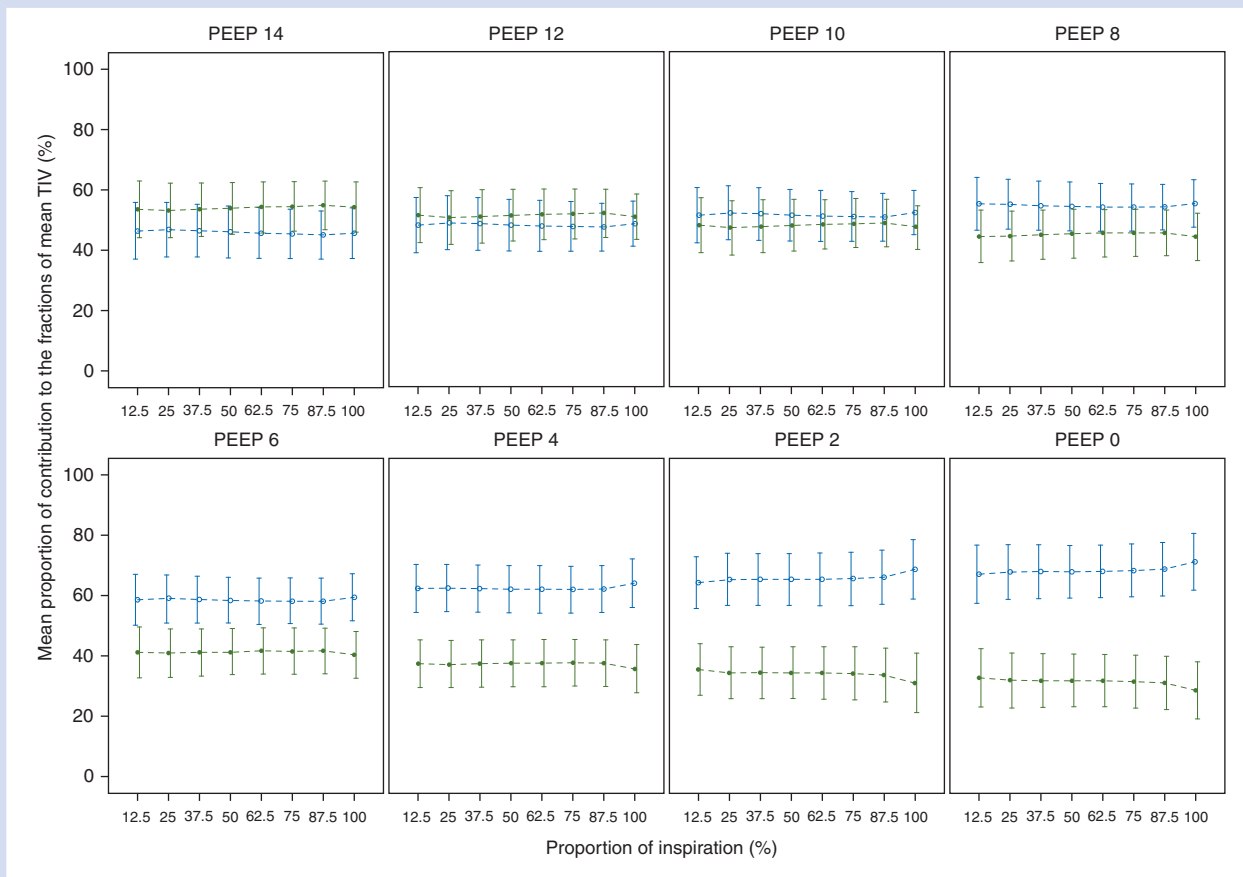
homogeneous ventilation as measured by EIT. In contrast, VD<sub>alv</sub>/VT<sub>alv</sub> and S<sub>III</sub> were insensitive to detect lung inhomogeneity. In addition, we found that in relatively healthy lungs, without the application of a RM, higher PEEP levels mainly induce airway distention rather than improve alveolar ventilation.

Recently, we demonstrated that homogeneous ventilation by means of ITV agreed well with a PEEP level, with the highest dynamic compliance.<sup>20</sup> We compared the whole dependent

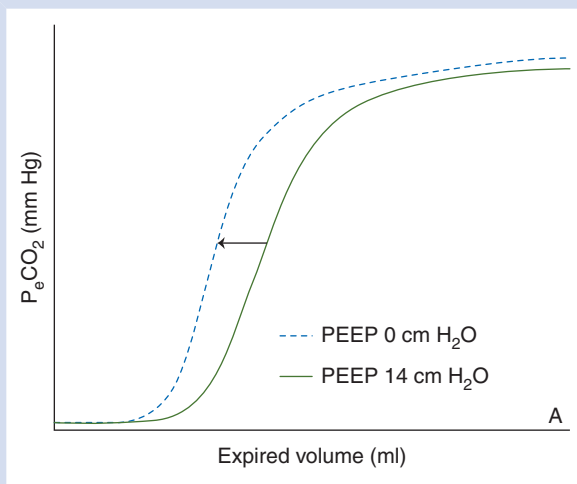
with the non-dependent lung region and found that the dependent lung region had the highest TIV at 12 cm H<sub>2</sub>O PEEP and decreased with each decremental PEEP step, indicating collapse in that region. At a PEEP level of 12 cm H<sub>2</sub>O the dependent and non-dependent lung regions were in balance. Above this PEEP the dependent region received most of the tidal volume, whereas below this PEEP the non-dependent region was predominantly ventilated. Thus EIT is the first device able to visualize collapse and hyperinflation of lung tissue at the bedside.

In an experimental study by Tusman and colleagues<sup>32</sup> a RM followed by decremental PEEP trial from 24 to 0 cm H<sub>2</sub>O in steps of 2 cm H<sub>2</sub>O was performed, in eight lung lavaged pigs. They investigated the ability of Vcap to detect optimal PEEP, as compared with computed tomography scans (CT), and found that dynamic compliance, VD<sub>alv</sub>/VT<sub>alv</sub> and Pa-ETCO<sub>2</sub> correlated best with the amount of non-aerated lung tissue as detected by CT-scans. Yang and colleagues<sup>33</sup> performed a decremental PEEP trial on eight lung-lavaged piglets. After a RM they stepwisely reduced the PEEP from 20 cm H<sub>2</sub>O to 4 cm H<sub>2</sub>O in steps of 4 cm H<sub>2</sub>O. At PEEP levels below 16 cm H<sub>2</sub>O they found that VD<sub>alv</sub>/VT<sub>alv</sub> was well correlated to non-aerated and normally aerated lung tissue as assessed by CT-scans. In addition, they found that the lowest S<sub>III</sub> value was closely related to best lung function. They concluded that VD<sub>alv</sub>/VT<sub>alv</sub> and S<sub>III</sub> are best predictors for alveolar collapse or hyperinflation. Therefore, from these two studies it can be concluded that Vcap is a reliable tool to detect both collapse and hyperinflation of lung tissue, as confirmed by CT-scans, which is the golden standard to assess alveolar collapse or hyperinflation.

Maisch and colleagues<sup>34</sup> performed an incremental and decremental PEEP trial in 20 anesthetized patients with healthy lungs, undergoing faciomaxillary surgery. During the PEEP trial, PEEP levels between 0 and 15 cm H<sub>2</sub>O were applied in PEEP steps of 5 cm H<sub>2</sub>O. They showed that after a RM physiological dead space was the lowest at 10 cm H<sub>2</sub>O PEEP, whereas static compliance of the respiratory system was the highest at that PEEP level. In addition, functional residual capacity and PaO<sub>2</sub> were insensitive parameters for detection of alveolar hyperinflation as



**Fig 3** Mean intratidal gas distribution (ITV) curve of all the patients at each PEEP step. Data are shown as mean (sd). ITV curve represents the mean percentile contribution (%) of ventilation distribution in non-dependent and dependent lung regions during the entire inspiration. At PEEP 12 and 10 cm H<sub>2</sub>O, intratidal gas distribution curves stayed close to each other, showing equal distribution to both regions. Dashed lines represents the interpolation lines; open circles=non-dependent region; solid circles=dependent region.



**Fig 4** Effect of increased PEEP on the Vcap curve. Figure 4 demonstrates the effect of two PEEP levels on the Vcap curve. As a result of the PEEP application the Vcap curve shifts to the right, indicating an increase in airway dead space.  $P_e\text{CO}_2$  = Expiratory carbondioxide pressure;  $V_{\text{cap}}$  = volumetric capnography.

they decreased with each PEEP step. Therefore, they concluded that physiological dead space and static compliance of the respiratory system, are suitable parameters for PEEP titration. Interestingly, we found that the  $VD_{\text{Bohr}}/VT$  and  $VD_{\text{Engelhoff}}/VT$  significantly decreased (Table 2), suggesting a reduction in V/Q mismatch, agreed well with the PEEP level at which the inspired air was homogeneously distributed to the dependent and non-dependent lung regions, as measured by EIT (Figs 2 and 3). However, at lower PEEP levels,  $VD_{\text{Bohr}}/VT$  and  $VD_{\text{Engelhoff}}/VT$  keeps decreasing indicating that this parameter adds no additional information about ventilation homogeneity at lower PEEP. In addition, the exhaled CO<sub>2</sub> per breath ( $VCO_{2,\text{br}}$ ) and the  $Sn_{\text{III}}$  both showed that elimination of CO<sub>2</sub> improved at lower PEEP levels (Table 2), but no optimum could be found. Therefore, in the present study design and study population, Vcap is not able to detect homogeneous ventilation.

Our finding in Vcap that PEEP mainly increases the airway-volume rather than improves alveolar ventilation is apparently in contradiction with the results of other studies published earlier.<sup>14 32 33</sup> Lower PEEP levels cause a shift of the Vcap curve to the left (Fig. 4A), indicating increased airway dead space at higher PEEP levels. These opposing results could be explained by the fact that post-cardiac surgery patients have relatively healthy lungs. Although the patients in our study may have atelectasis,

there is no question of surfactant depletion and we assume that the atelectasis is recruited without the need for high PEEP levels. In contrast, the previous studies used the lung-lavaged models with large amount of shunt area, which are recruitable at high PEEP levels. However, Nieman and colleagues<sup>35–41</sup> found during *in vivo* microscopy in normal and surfactant deactivated pigs, during ventilation with tidal volumes of 6, 12 and 15 ml kg<sup>-1</sup>, that in normal lungs the alveolar area at end-inspiration and end-expiration (I-E<sub>A</sub>) did not differ between the three different tidal volumes, whereas in the surfactant deactivated lungs I-E<sub>A</sub> increased significantly at higher tidal volumes. In a next study<sup>35</sup> they used different tidal volumes (6, 12, 15 ml kg<sup>-1</sup>) and PEEP levels (5, 10, 20 cm H<sub>2</sub>O) and found that I-E<sub>A</sub> did not change in normal lungs, whereas in surfactant deactivated lungs the alveolar size significantly increased, as compared with normal lungs with equal settings. From these studies it can be concluded that in normal lungs with stable alveoli, PEEP volume results in airway distention, whereas in ALI/ARDS lungs, PEEP keeps the alveoli open and stabilized. In addition, airway distention as a result of high PEEP might also explain why physiological dead space calculated according to Enghoff, increases at higher PEEP levels, whereas PaCO<sub>2</sub> did not significantly differ during the different PEEP levels, except at ZEEP. This explains the absence of a change in SIII and the left shift of SII in the Vcap curves at decreasing PEEP levels in our patients.

There are limitations to this study. Firstly, we did not apply a RM before the PEEP reduction, in order to avoid serious deterioration of haemodynamics in the direct postoperative period. It has been shown that Vcap parameters after an RM are more sensitive to detect changes in CO<sub>2</sub> elimination, as compared with PEEP trials without an RM.<sup>42–43</sup> Secondly, EIT describes the ventilation distribution in a lung slice of approximately 5–10 cm thick.<sup>42–44</sup> Therefore, the image obtained from the single cross-section does not represent the entire lung. In addition, PEEP increases results in changed chest size and intrathoracic blood volume. However, by using two large regions of interest, the corresponding lung tissue to the individual pixel is limited. The effect of changed intrathoracic blood volume is limited, as EIT is a hundred times more sensitive to impedance changes caused by aeration, as compared with impedance changes caused by the cardiac system.<sup>45–47</sup> Thirdly, the time intervals of 5–10 min at each PEEP level might be too short to achieve a steady state for obtaining reliable data. However, a recent study showed that elimination of CO<sub>2</sub> reached a new stable period within five min of a PEEP change.<sup>48</sup> It is known that Vcap is influenced by both the cardiac system and the pulmonary system. In this study we did not perform cardiac output measurements. Therefore we cannot assess Vcap changes induced by reduced cardiac output at higher PEEP levels. However, we found no significant differences in bp and heart rate between PEEP 14 and ZEEP. Therefore, we assume that the influence of the change in cardiac output with different PEEP levels on dead space was not profound.

## Conclusion

In postoperative cardiac patients with low shunt fractions, dead space calculations according to Bohr and Enghoff agreed well with EIT. EIT is able to detect the optimal PEEP level for equal distribution of inspired air, among the non-dependent and the dependent regions of the lungs. In contrast, VD<sub>alv</sub>/VT<sub>alv</sub> and Sn<sub>III</sub> were unable to detect this distribution. In addition, we found that in relatively healthy lungs, applying higher PEEP levels mainly induced airway distention rather than improvement of alveolar ventilation.

## Authors' contributions

Study design/planning: P.B., D.G.  
Study conduct: P.B., A.S., B.H., T.W.  
Data analysis: P.B., A.S., B.H., T.W., D.H., D.G.  
Writing paper: P.B., A.S., B.H., T.W., D.H., D.G.  
Revising paper: all authors

## Supplementary material

Supplementary material is available at *British Journal of Anaesthesia* online.

## Declaration of interest

None declared.

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