Transpulmonary thermodilution cardiac output measurement is not affected by severe pulmonary oedema: a newborn animal study

A. Nusmeier1*, S. Vrancken2, W. P. de Boode2, J. G. van der Hoeven1 and J. Lemson1

1 Department of Intensive Care Medicine and 2 Department of Pediatrics, Radboud University Nijmegen Medical Centre, Nijmegen, The Netherlands
* Corresponding author. E-mail: a.nusmeier@ic.umcn.nl

Editor’s key points
- The effect of pulmonary oedema on reliability of transpulmonary thermodilution cardiac output measurement was investigated.
- Cardiac output was measured in newborn lambs in the presence of increased extravascular lung water.
- Measurement of cardiac output by transpulmonary thermodilution is not affected by severe pulmonary oedema in a newborn lamb model.

Background. The transpulmonary thermodilution (TPTD) technique is widely used in clinical practice for measuring cardiac output (CO). This study was designed to investigate the influence of various levels of pulmonary oedema on the reliability of CO measurements by the TPTD method.

Methods. In 11 newborn lambs pulmonary oedema was induced using a surfactant washout technique. Serial CO measurements using TPTD (CO_{TPTD}) were performed at various amounts of lung water. Simultaneously, CO was measured by an ultrasound flow probe around the main pulmonary artery (CO_{MPA}) and used as the standard reference. CO was divided by the body surface area to calculate cardiac index (CI). Data were analysed using correlational statistics and Bland–Altman analysis.

Results. One lamb died prematurely. A total of 56 measurements in 10 lambs were analysed with a median CI_{MPA} of 2.95 (IQR 1.04) litre min^{-1} m^{-2}. Mean percentage increase in extravascular lung water (EVLW) between the start and the end of the study was 126.4% (SD 40.4). Comparison of the two CO methods showed a mean bias CI of −0.16 litre min^{-1} m^{-2} (limits of agreement ± 0.73 litre min^{-1} m^{-2}) and a percentage error of 23.8%. Intraclass correlation coefficients were 0.91 (95% CI 0.81–0.95) for absolute agreement and 0.92 (95% CI 0.87–0.95) for consistency. Acceptable agreement was confirmed by a tolerability-agreement ratio of 0.39. The within-subject correlation between the amount of EVLWI and the bias between the two methods was not significant (r=0.02; P=0.91).

Conclusions. CO measurements by the transpulmonary thermodilution technique over a wide range of CI values are not affected by the presence of high EVLWI. The slight underestimation of the CO is independent of the amount of pulmonary oedema.

Keywords: cardiac output; children; haemodynamic; monitoring; pulmonary oedema; thermodilution

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in adults analysed the dependency of CO measurements on the amount of lung oedema. However, as infants and young children have a much higher EVLW indexed to body weight (EVLWI) than adults, the impact of these increased amounts of lung oedema may be more pronounced. Paediatric studies are, to our knowledge, not available. Therefore, this newborn animal experiment was designed to investigate the influence of various levels of pulmonary oedema on the reliability of CO measurements.

Methods

General

This experiment was performed in accordance with Dutch legislation concerning guidelines for the care and use of laboratory animals and was approved by the local ethics committee on animal research of the Radboud University Nijmegen Medical Centre (RUNMC Licence number RU-DEC 2010-034; CDL-project number 33078). Eleven lambs were studied under general anaesthesia. Premedication consisted of the i.m. administration of midazolam (0.2 mg kg$^{-1}$), ketamine (10 mg kg$^{-1}$), and i.v. administration of propofol (2 mg kg$^{-1}$). General anaesthesia was maintained using inhalation of isoflurane (1–1.5 vol%) and the continuous i.v. administration of sufentanil (20 μg kg$^{-1}$ h$^{-1}$), midazolam (0.2 mg kg$^{-1}$ h$^{-1}$), ketamine (10 mg kg$^{-1}$ h$^{-1}$), and pancuronium (0.02 mg kg$^{-1}$ h$^{-1}$) after a loading dose of 0.05 mg kg$^{-1}$. The depth of anaesthesia was repeatedly assessed by painful stimuli and clinical parameters such as heart rate, spontaneous ventilation, and elevated arterial pressure. The depth of anaesthesia was adjusted when necessary. During the experiment, continuous i.v. dextrose 10% 2 ml kg$^{-1}$ h$^{-1}$ was administered. The lambs were intubated orotracheally using a 4–6 mm (inner diameter) cuffed tracheal tube (Kruse, Marslev, Denmark). The lungs were mechanically ventilated in a pressure-controlled mode using tidal volumes of ~10 ml kg$^{-1}$ (Datex-Ohmeda anaesthesia machine) and an inspiratory-to-expiratory ratio of 1:2. Normocapnia, guided by capnography with the CO$_2$SMO Plus Respiratory Profile Monitor (Model 8100, Respironics, Pittsburgh, USA), was achieved by adjusting the minute volume ventilation to maintain an end-tidal CO$_2$ tension between 4.0 and 5.5 kPa. Impaired oxygenation was treated by adjusting the positive end expiratory pressure (PEEP) and the fraction of inspired oxygen ($F_{IO_2}$) to maintain the oxygen saturation >95%. A servo-controlled heating mattress and an external heating lamp were used to maintain core temperature between 38 and 40°C. At the end of the experiment, the animals were killed with an overdose of pentobarbital (150 mg kg$^{-1}$ i.v.).

Instrumentation

Immediately after induction of anaesthesia, a thermal-dye-dilution probe (PV2023, Pulsion, Germany) equipped with a thermistor for the detection of changes in blood temperature and a fibreoptic probe to detect plasma levels of green dye was inserted in the femoral artery. In the contralateral femoral vein, a central venous catheter (5Fr, 2 lumen, 13 cm, Arrow, Germany) was inserted for the administration of fluid and drugs. At the same site a femoral artery catheter (20 Ga, single lumen, 12 cm, Arrow, Germany) was introduced for arterial pressure monitoring and blood sampling. All intravascular catheters were inserted by a surgical cut-down technique. A left-sided thoracotomy was performed and the remains of a native ductus arteriosus were ligated. An ultrasound transit time perivascular flow probe (10 or 12 mm) (PAX series, Transonic Systems, Ithaca, NY) was placed around the main pulmonary artery to measure reference CO (CO$_{MPA}$). The flow probe signal was checked for zero flow values directly postmortem. Ultrasound transit time flow probes use a two-way ultrasound technique. By calculating the difference between transit times upstream and downstream, the blood flow (Q$_{MPA}$) is measured. Care was taken to avoid air within the flow probe by applying sufficient quantities of acoustic gel. After the placement of the flow probe, the thorax was closed. The animals were positioned either supine or lying on the right side throughout the experiment.

Pulmonary oedema was induced using a surfactant washout lavage model. In short, lambs underwent repetitive saline lavages (10–35 ml kg$^{-1}$ lavage$^{-1}$ 37°C NaCl 0.9%) of the lung in order to induce surfactant depletion and provoke acute lung injury (ALI). Before the lavages the lambs were pre-oxygenated using an $F_{IO_2}$ of 1.0. After the lavages, the animals were stabilized for 30 min before measurements of ventilatory and haemodynamic parameters and blood gases were obtained. Between lavages the PEEP and minute volume ventilation were increased to maintain oxygen saturation and end-tidal CO$_2$ within the normal range.

Transpulmonary thermodilution

Transpulmonary thermodilution CO (CO$_{TPTD}$) was measured by injection of 5 ml ice-cold saline (NaCl 0.9%) into the femoral venous catheter. Changes in temperature were detected by the thermistor connected to a COLD monitor (Pulsion, Munich, Germany). The theoretical background of measuring CO by analysis of the dilution curves and calculation using the Steward Hamilton equation is described elsewhere. Besides CO, blood volumes and extravascular lung water can be calculated from the measurement of the mean transit time (MTTs) and downslope time (Dst) of the dilution curves. Before a series of thermodilution measurements, the central venous catheter was flushed with 1–2 ml of ice-cold saline. Each thermodilution curve was visually inspected for artifacts or signs of an inadequate measurement. We used the mean value of three bolus injections of 5 ml of ice-cold (<10°C) saline.

Other measurements

We measured invasive arterial pressure and central venous pressure, continuous electrocardiogram, heart rate, arterial oxygen saturation, end-tidal CO$_2$, respiratory frequency, tidal volume, airway pressures, and body core temperature. During the thermodilution measurements, all other
haemodynamic variables, including CO_{CITPTD}, were recorded simultaneously with a 200-Hz sampling rate using a computer system with special biomedical registration software (Poly, Inspektor Research Systems, Amsterdam, The Netherlands). The exact time span of the dilution measurement was marked in the registration system. The reference CO was calculated using the mean value of CO_{CITPTD} measurements over the same three periods as the mean value of three consecutive TPTD measurements.

**Protocol**

After instrumentation, baseline measurements of CO (TPTD and CO_{CITPTD}) respiratory and haemodynamic parameters and blood gases were obtained. Repetitive saline lavage procedures were performed in either one or two subsequent sessions of 10–30 ml kg⁻¹, depending on the recovery of the lambs during the procedure. After each lavage procedure, a pause of 30 min was instituted for cardiorespiratory stabilization, followed by repeated measurements of the above-mentioned parameters. A blood transfusion was administered if the haemoglobin (Hb) was < 3.5 mmol litre⁻¹. Dobutamine or epinephrine was administered when indicated.

**Statistical analysis**

The CO values were indexed (CI) to body surface area (BSA). The BSA of the lambs was calculated using the following formula: BSA = weight²/³ × 0.121. The statistical variability in the CO during the measurement periods is expressed as the coefficient of variance (CV). This is calculated by dividing the standard deviation (SD) of the mean by the mean of the CO_{CITPTD} results during each separate measurement period. The results are expressed as percentages (×100% × SD/ mean). We defined a CV of ≤5% as acceptable for reliable CO_{TPTD} measurements. Intraclass correlation coefficients, using the two-way mixed model, were calculated for consistency and absolute agreement between the CO measurement methods. In addition, data were analysed using the method described by Bland and Altman. The difference between the two methods (bias) was calculated by subtracting the value of CO_{CITPTD} from CO_{TPTD}. The bias was plotted against the mean CI (CI_{TPTD} + CI_{CITPTD})/2. The limits of agreement (LOA) were calculated by multiplying the SD of the bias by 1.96. The percentage error was calculated using the following formula: [(1.96 × SD of the bias)/mean CI_{CITPTD}] × 100%, using mean CI_{CITPTD} as the reference. As the number of measurements per animal varied, we corrected for the repeated measurements. The strength of agreement was also calculated by the agreement-tolerability-interval ratio, with acceptable agreement defined as a ratio < 1. The tolerability interval is estimated from the 95% range of the observed CI data. The agreement interval is calculated from the range of the agreement-tolerability-interval ratio, with acceptable agreement-tolerability-interval ratio was 0.39.

Separate analysis of the lower and higher CI measurements showed comparable percentage errors 22% ([CI<3.0 litre min⁻¹ m⁻²]; n=28; mean bias −0.06 (LOA 0.51) litre min⁻¹ m⁻²) and 23% ([CI≥3.0 litre min⁻¹ m⁻²]; n=28; mean bias −0.27 (LOA 0.85) litre min⁻¹ m⁻²). We subsequently determined the bias and level of agreement of the CI measurements for different amounts of EVLWI and showed a mean bias of −0.28 (LOA 0.57) litre min⁻¹ m⁻² and percentage error of 16.1% in the lowest quartile of EVLWI vs −0.27 (LOA 0.38) litre min⁻¹ m⁻² and percentage error of 21.8% in the highest quartile of EVLWI. Differences in the bias and LOA were significant in the low/high CI (P=0.014 and P=0.008) groups, but not in the low/high EVLWI (P=0.75 and P=0.84) groups.

The initial median EVLWI was 16.0 ml kg⁻¹ (IQR 3.8), which increased after multiple lung lavage procedures to a final median EVLWI of 36.3 ml kg⁻¹ (IQR 9.1). The mean percentage increase of EVLWI between the start and the end of the study was 126.4% (SD 40.4). The increment of EVLWI worsened oxygenation. The median PAo₂/FIO₂ ratio decreased from 403 (IQR 173) to 73 (IQR 54). The median PEEP had to be increased during the experiment from 5 cm H₂O (IQR...
0) up to 16 cm H₂O (IQR 10). The differences in these parameters between the start and final measurements were analysed by the Wilcoxon signed-rank test (all \( P = 0.02 \)) and are shown in Figure 2. The overall COTPTD measurements underestimated slightly the CO with 6%, increasing up to 7% bias in the highest \( \frac{Cl_{TPTD} + Cl_{MPA}}{2} \) EVLWI values.

When we compared the bias and LOA between the two CO measurements during low PEEP \( (n=22) \) and high PEEP \( (n=32) \), these differences were not statistically significant \( (P=0.98 \text{ and } P=0.11) \). The within-subject correlations between the amount of EVLWI, \( PaO_2/FIO_2 \) ratio, or the PEEP and the bias between the two CO measurement methods were \( r = -0.02 \) \( (P=0.91) \), \( r = 0.09 \) \( (P=0.56) \) and \( r = 0.10 \) \( (P=0.5) \), respectively, and are illustrated in Figure 3.

## Discussion

In this newborn animal model, the TPTD method accurately measured CO in the presence of severe pulmonary oedema (increase more than 125%). The bias and LOA were small and the percentage error around 24% is within the range of acceptance. \(^{22} \) The high intraclass correlation coefficient and the low (<0.5) agreement-tolerability-interval ratio, taking the reference range of the observed data of the study as tolerability interval into account, support the acceptable strength of agreement and imply the EVLWI to be irrelevant in this study. \(^{24} \) The initial EVLWI values in our paediatric animal model were higher than normal indexed EVLW values known from adult studies, but in agreement with the high EVLWI found in young children. \(^{14} - ^{16} \) \(^{26} \) Our results show no important dependency of the bias between the two CO methods and the amount of EVLW, but rather a scatter of differences that need to be considered.

It has been estimated that with the TPTD method, up to 9% of the thermal indicator may be lost during passage through the pulmonary circulation. \(^ {27} \) In critically ill paediatric patients without pulmonary oedema, TPTD overestimates the CO up to 4.4% compared with pulmonary artery thermodilution. \(^6 \) This is in agreement with adult studies. \(^6 - ^{10} \) \(^ {28} \) Our results did not show overestimation of the CO\(_{TPTD}\). An explanation for this discrepancy could be that, in contrast to others, we used an ultrasound flow probe as reference method, which is not influenced by other factors. Most human studies use pulmonary artery thermodilution as the reference method, which is influenced by the traversing distance of the indicator and the transient effect of ice water on the heart rate. \(^{29} \) \(^ {30} \)

Only a few adult studies focus on the influence of pulmonary oedema on the reliability of CO measurements using the TPTD technique. A retrospective study in adult surgical intensive care patients showed no dependency of the bias between TPTD and pulmonary artery thermodilution CO measurements on the amount of pulmonary oedema. \(^ {12} \) In this study, however, the amount of EVLWI was relatively low (mean 9.1 ml kg\(^{-1}\)) compared with the much higher EVLWI values in children. Another adult study in ARDS patients with a high mean EVLWI of 20.2 ml kg\(^{-1}\) showed similar results. \(^ {13} \) This may be explained by re-entry of the lost cold thermal indicator into the flowing blood. An older

### Table 1 Characteristics of the lambs

<table>
<thead>
<tr>
<th>Lamb</th>
<th>Weight (kg)</th>
<th>Age (days)</th>
<th>Total lavages (ml kg(^{-1}))</th>
<th>CO measurements</th>
<th>Mean ( CO_{MPA} (\pm \text{so}) ) (litre min(^{-1}))</th>
<th>Mean ( CI_{MPA} (\pm \text{so}) ) (litre min(^{-1}) m(^{-2}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>8.1</td>
<td>21</td>
<td>125</td>
<td>7</td>
<td>1.89 (0.13)</td>
<td>3.86 (0.27)</td>
</tr>
<tr>
<td>2</td>
<td>7.4</td>
<td>21</td>
<td>303</td>
<td>5</td>
<td>1.06 (0.29)</td>
<td>2.31 (0.63)</td>
</tr>
<tr>
<td>3</td>
<td>4.1</td>
<td>7</td>
<td>293</td>
<td>6</td>
<td>1.00 (0.09)</td>
<td>3.24 (0.29)</td>
</tr>
<tr>
<td>4</td>
<td>7.4</td>
<td>14</td>
<td>68</td>
<td>2</td>
<td>2.48 (0.81)</td>
<td>5.39 (1.77)</td>
</tr>
<tr>
<td>5</td>
<td>10.2</td>
<td>17</td>
<td>294</td>
<td>6</td>
<td>2.06 (0.49)</td>
<td>3.62 (0.86)</td>
</tr>
<tr>
<td>6</td>
<td>9.4</td>
<td>18</td>
<td>191</td>
<td>4</td>
<td>2.05 (0.14)</td>
<td>3.81 (0.26)</td>
</tr>
<tr>
<td>7</td>
<td>Died</td>
<td></td>
<td></td>
<td></td>
<td>Excluded</td>
<td>Excluded</td>
</tr>
<tr>
<td>8</td>
<td>8</td>
<td>15</td>
<td>420</td>
<td>8</td>
<td>1.01 (0.26)</td>
<td>2.08 (0.54)</td>
</tr>
<tr>
<td>9</td>
<td>9.9</td>
<td>16</td>
<td>242</td>
<td>3</td>
<td>1.61 (0.25)</td>
<td>2.89 (0.45)</td>
</tr>
<tr>
<td>10</td>
<td>11.5</td>
<td>19</td>
<td>184</td>
<td>7</td>
<td>1.82 (0.25)</td>
<td>2.95 (0.40)</td>
</tr>
<tr>
<td>11</td>
<td>12.3</td>
<td>22</td>
<td>98</td>
<td>8</td>
<td>1.61 (0.23)</td>
<td>2.50 (0.36)</td>
</tr>
</tbody>
</table>

Fig 1 Bland–Altman plot comparing the cardiac index values by the TPTD method \((CI_{TPTD})\) and the peri-vascular flow probe around the main pulmonary artery \((CI_{MPA})\).
study found a decrease in thermal loss as EVLW accumulates with a negative correlation between the bias and the EVLW. The measured values of EVLWI in this adult study ranged from 1.9 up to 27.5 ml kg\(^{-1}\). In our study, the EVLWI values ranged from 12.8 up to 60.2 ml kg\(^{-1}\). In agreement with the latter study, our results also show a slight (6%) underestimation of the mean COTPTD.

These findings suggest that in the presence of severe pulmonary oedema, other factors compensate for the possible loss of the thermal indicator. Factors governing the temperature exchange between blood and interstitial fluid are the velocity of diffusion of the cold indicator, the surface area for exchange, the volume of the interstitial fluid into which cold diffuses, and the flow of blood, which determines the time during which temperature exchange must occur. So, as a result of perfusion alterations induced by pulmonary oedema, PEEP or both, the surface area for exchange is reduced compensating thermal indicator loss in oedematous lung tissue. However, our study could not demonstrate an effect of PEEP on the bias between CO measurements. Blood flow alteration is another factor influencing the loss of thermal indicator. High CO states may not allow sufficient time for equilibration.
with the extra vascular fluids and therefore less thermal indicator may be lost.9 On the other hand, low CO values result in more indicator loss at lower flow rates.32 33 This influence of blood flow on bias is confirmed in our study comparing low with high CO values. However, the error margin of the measurements remains fractionally the same. The higher the CO, the wider the LOA and vice versa, resulting in comparable percentage errors at high and low CO values.

**Limitations**

Despite variable haemodynamic circumstances, we included only measurements during stable blood flow. We corrected for the unequal and repeated CO measurements per lamb. The animals were either lying supine or on their right side, which could influence the distribution of pulmonary oedema, atelectases, and ventilation-perfusion mismatch and may have influenced our measurements in the comparison between the animals. We had no clinical indication to assume intracardiac shunting. We used a common model to induce EVLW increment with well-described pathophysiological and morphological characteristics.17 34 35 However, this surfactant depletion model does not share all features of ALI/ARDS in humans. Finally, the difference in normal values of EVLWI between children and adults may not be real but the result of a difference in age-related changes in the ratio of lung weight to body weight ratio.36

**Conclusions**

Haemodynamic monitoring plays a crucial role in the treatment of critically ill patients. In particular, patients with capillary leakage and ALI require tight fluid management avoiding the risk of overzealous fluid administration while maintaining sufficient intravascular volume status. The transpulmonary thermodilution technique provides reliable CO monitoring over a wide range of clinical conditions. Our study shows that CO measurements are not affected by severe pulmonary oedema in a newborn lamb model.

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**Declaration of interest**

None declared.

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