Respiratory modulations in the photoplethysmogram (DPOP) as a measure of respiratory effort

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Abstract  DPOP is a measure of the strength of respiratory modulations present in the pulse oximetry photoplethysmogram (pleth) waveform. It has been proposed as a non-invasive parameter for the prediction of the response to volume expansion in hypovolemic patients. The effect of resistive breathing on the DPOP parameter was studied to determine whether it may have an adjunct use as a measure of respiratory effort. Healthy volunteers were tasked to breathe at fixed respiratory rates over a range of airway resistances generated by a flow resistor inserted within a mouthpiece. Changes in respiratory efforts, effected by the subjects and measured as airway pressures at the mouth, were compared to DPOP values derived from a finger pulse oximeter probe. It was found that the increased effort to breathe at fixed respiratory rates over a range of airway resistances generated by a flow resistor inserted within a mouthpiece, were compared to DPOP values derived from a finger pulse oximeter probe. It was found that the increased effort to breathe manifests itself as an associated increase in DPOP. Further, a relationship between DPOP and percent modulation of the pleth waveform was observed. A version of the DPOP algorithm that corrects for low perfusion was implemented which resulted in an improved relationship between DPOP and PPV. Although a limited cohort of seven volunteers was used, the results suggest that DPOP may be useful as a respiratory effort parameter, given that the fluid level of the patient is maintained at a constant level over the period of analysis.

Keywords  Hemodynamic monitoring · Fluid responsiveness · Pulse oximetry · DPOP · PPV

1 Introduction

Respiratory effort is a context specific term defined relative to the clinical setting. A range of physiological measurements may be used in its assessment including maximal inspiratory and expiratory pressures [1]; pleural pressures, esophageal pressures and work of breathing [2]; changes in nasal cannula pressures [3]; changes in blood pressures or pulse transit times [4, 5]; changes in EMG signals [6]; chest band extensions [7]; visual observation of signs of distress and muscle use [8]; changes in coloration [9]; and respiratory sounds [10]. It is now recognised that respiratory components clearly manifest in the photoplethysmogram (pleth) across a range of conditions [11–13]. Analysis of the frequency of these components has led to the development of respiratory rate (RR) algorithms for incorporation within pulse oximeter devices [14, 15]. However, the strength of these modulations also provides information on the thoracic pressure changes, associated with the effort to breathe. Respiratory effort manifests in the pleth waveform as a variety of distinct modulations which has led to the development of various technologies for its measurement [16–19].

This present study explored the potential for using the pulse oximetry photoplethysmographic fluid responsiveness parameter, DPOP, as a measure of respiratory effort. DPOP is a non-invasive measure of the normalized strength of respiratory modulations present in the pleth waveform [20]. It is defined as [21]:

$$DPOP = \frac{(AMP_{\text{max}} - AMP_{\text{min}})}{(AMP_{\text{max}} + AMP_{\text{min}})/2}$$

(1)

where $AMP_{\text{max}}$ and $AMP_{\text{min}}$ are the maximum and minimum amplitudes of the cardiac pulse waveforms in the pleth during a respiratory cycle. These are illustrated in
Fig. 1, where the cardiac pulse component of the pleth is shown being modulated by respiratory activity. DPOP has been proposed as a measure of the response to fluid administration in mechanically ventilated patients. As such it represents a non-invasive alternative for pulse pressure variation (PPV) [22] with many studies showing favourable correlation between the two parameters [23–29]. DPOP is derived from the change (or delta) in the pulse oximetry photoplethysmogram (POP) waveform.

In this study, data from healthy volunteers undergoing coached breathing in a laboratory setting was employed to study the effect of respiratory effort on DPOP. The DPOP parameter was compared with PPV, derived from the arterial blood pressure waveform which may also be used in practice to determine fluid responsiveness. The effect of low perfusion on the DPOP parameter was also considered where the performance of a modified DPOP parameter, corrected for low perfusion, was studied. This low perfusion correction is fully documented elsewhere [30, 31].

2 Methods

2.1 Protocol

With institutional review board approval and written informed consent, seven healthy adult volunteers were enrolled at the Covidien RMS clinical laboratory in Boulder, CO. These volunteers had been pre-screened and completed physical exams. During the trials, the subjects were positioned comfortably in a chair with the pulse oximeter sensor attached (Max-A, Covidien, Boulder, CO). A facemask was attached to each subject consisting of a spirometer containing a number of interchangeable flow resistors [5, 20 and 50 cmH2O/l/s linear resistors (Hans Rudolf Inc., Kansas City, MO)]. The blood pressure signal from an intra-arterial line was also recorded. A synchronized acquisition of the pulse oximeter, arterial pressure and airway pressure signals was performed via a custom data acquisition system during the whole procedure and saved to a laptop for later analysis.

The subjects were asked to breathe at a fixed RR of 8 breaths-per-minute (brpm) through the spirometer. This was facilitated by following a metronome displayed on a computer screen next to the subject. Each breathing task was maintained for a 1-min period separated in time by a 2-min recovery period in order to allow the subject to relax and breathe freely in preparation for the next 1-min task.

Each subject was initially asked to breathe through the selected flow resistor at a consistent effort for a 1-min period at what they felt was an appropriate effort for that resistance. This is labelled (M). For the second 1-min effort task, the subject was asked to breathe with more effort through the resistor. This is labelled (H). During the third task the subject was asked to breathe with less effort through the resistor than the initial effort (S), and during the fourth task the subject was asked to breathe in a similar manner to the first attempt (M). The medium (M) efforts allowed the patient to familiarize him/herself with the apparatus and to set a baseline. The maximum (H) and minimum (L) effort breathing values were then extracted for analysis. This provided two 1-min segments of pleth and pressure signals at either extreme of air pressures at the mouthpiece. The equipment also allowed for the option to set the flow resistance only on inhalation or only on exhalation. The regimes undertaken by the seven subjects are summarised in Table 1 where it can be seen that five subjects encountered flow resistance on inhalation and two on exhalation. Each series of tasks was repeated for the three flow resistances—5, 20 and 50 cmH2O/l/s. Two subjects had difficulty breathing at the highest flow resistor level of 50 cmH2O/l/s and did not complete this task.

2.2 Data analysis

An example of the acquired signals is shown in Fig. 2a from one of the task sets for a linear resistor setting of 20 cmH2O on exhalation only. The sequential medium, high, low and medium airflow efforts are marked on the plot (M, H, L, M). During these periods the subject breathed at 8 brpm and outside these periods the subject was free to relax and breathe freely at his or her natural rate. Figure 2b shows a zoom of the area marked by an arrow in Fig. 1 where large amplitude modulations can be observed in the pleth signal. It is clear in Fig. 2b that these modulations are also present in the blood pressure signal. DPOP was calculated from the pleth using the algorithmic method described in [30] but with shorter averaging employed as the signal segments were only of 1 min duration (i.e. here DPOP was only averaged over the 1 min window). PPV was computed using the same formula as DPOP. The corresponding air pressure modulation strength (APM) was calculated as the mean of the breath-by-breath maxima of the air pressure modulations. Plots of DPOP versus PPV, DPOP versus APM and PPV versus APM were generated, together with corresponding difference plots.
It can also be observed from the lower plot of Fig. 2b that the respiratory modulations between 380 and 420 s rise up on one of a number of slower waveforms that can be seen in the signal segment. Note that such longer-period modulations are very typical of raw photoplethysmographic signals and are due to a variety of causes:

Table 1 The available subject data sets

| Patient ID | Resistor direction | C = 20 | C = 5 | C = 50 |
|------------|--------------------|--------|-------|--------|
| RE002      | Inhale             | Y      | Y     | N      |
| RE003      | Exhale             | Y      | Y     | Y      |
| RE005      | Inhale             | Y      | Y     | N      |
| RE006      | Exhale             | Y      | Y     | Y      |
| RE008      | Inhale             | Y      | Y     | Y      |
| RE009      | Inhale             | Y      | Y     | Y      |
| RE011      | Inhale             | Y      | Y     | Y      |

Flow resistor values ‘C' in cmH₂O/l/s

Y task completed, N task not completed

Fig. 2 Airflow (top), Arterial blood pressure (middle) and pleth (bottom). a Signals for the whole four tasks. b Zoom into the signal segments denoted by the arrow in (a)
vasomotion, blood pressure changes, slow/slight positional changes or movements, etc. Normally, all such signal information (including the respiratory modulations) is filtered out in standard pulse oximeter devices.

3 Results

The relationships between APM, PPV and DPOP for the data are shown in Fig. 3. The increasing effort for each task is indicated by linking each of the L–H data pairs by a line where the lower respiratory effort data point is indicated by the smaller symbol and the larger respiratory effort data point by the larger symbol. A least squares fit is shown in Fig. 3a between DPOP and PPV showing the positive correlation with $R = 0.533$.

Figure 4 contains the difference plots associated with the data in Fig. 3. These comprise the difference in the parameters APM, PPV and DPOP where the parameter values at low subject effort is subtracted from those at high subject effort (small and large symbols in Fig. 3). The plots allow us to see at a glance the general trending direction of

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**Fig. 3** Relationships between airway pressure, DPOP and PPV. a Relationship between DPOP and PPV. b Relationship between PPV and APM. c Relationship between DPOP and APM. Subjects are differentiated by color. Linear resistors of 5/20/50 cmH$_2$O/l are indicated by solid/dashed/dotted lines with circles/triangles/squares respectively. High/low breathing efforts are indicated by large/small symbols at the end of each line. All data were collected with a coached breathing at a respiration rate of 8 brpm.
the parameters as the subject switches from low effort breathing to high effort breathing. The parameters all appear to increase in magnitude with increasing respiratory effort with the exception of two outliers. The outlier in the lower-right hand quadrant corresponds to a signal acquisition glitch and the one just inside the top-left quadrant corresponds to a relatively low value of change in modulation strength and thus is dominated by signal noise.

The relationship between DPOP and the percent modulation (PMod) is shown in Fig. 5. PMod, sometimes also called perfusion index, is the ratio of the AC to DC component in the pleth signal. Figure 6 shows the updated version of Figs. 3a and 4a using an enhanced DPOP algorithm with a correction for PMod (described in [31]). It can be seen that the correlation coefficient between PPV and DPOP has improved from $R = 0.533$ to 0.739.

**4 Discussion**

Numerous attempts have been made over recent years to assess respiratory effort from the photoplethysmogram. This includes the work by Hartert et al. [16] who used the baseline modulation as a measure of airway obstruction in a group of patients with obstructive airway disease (asthma or COPD). They found that the measure gave good separation when used to identify those patients with abnormal pulsus paradoxus from those with physiologic pulsus (defined either side of a 10 mmHg threshold). Later, Arnold [17] found a relationship between the variation in waveform area and height of the cardiac pulse through the respiratory cycle and changes in mouth pressure in a study of healthy young adults who breathed against progressively increasing levels of mouth pressure. More recently Perel [19] suggested that pleth-based fluid responsiveness

![Fig. 4 Difference plots of respiratory effort induced parameter changes. a DPOP change versus PPV change; b PPV change versus mean absolute airway pressure change; c DPOP change versus mean absolute airway pressure change](image-url)
measures may be clinically important for the detection of respiratory effort increases due to upper airway obstructions. In fact, the Perel paper prompted the recent editorial by Cannesson and Shelley [32] who asked how the same pulse oximeter waveform could "monitor oxygen saturation, guide fluid-replacement therapy, and spot airway obstruction?" The answer to this lies in two parts. Firstly, oxygen saturation involves a different phenomenological aspect of the photoplethysmogram than the other two where signals from two wavelengths of light are employed and a ratio of normalized cardiac pulse amplitudes is used to derive the parameter. However, fluid responsiveness and the effects of airway obstruction require analysis of the same (respiratory) components in a single signal. The answer to how these can provide both fluid volume status and a respiratory effort measure lies in the dynamics of the patient physiology during the measurement. If breathing at a constant effort then the effect of differing fluid volume levels can be monitored. If however, volume status is held constant—as it is assumed for the subjects in the current study—the effect of changing respiratory efforts may be monitored.

The analysis compared DPOP to PPV. Although the pulsatile pleth and blood pressure waveforms resemble each other, these are not equivalent physiological measures. Nonlinear coupling exists between the pressure pulses and the vessel-wall mechanical activation causing the pulsed light change in the pleth. This is discussed in more detail elsewhere [31]. However, both contain both cardiac and respiratory modulations and both are currently used as fluid responsiveness parameters in practice. Another popular metric for assessing fluid responsiveness through the pleth is the pulse variability index (PVI) parameter. However, as discussed elsewhere by the author, [20] PVI is a proprietary algorithm of the Masimo Corporation (Masimo Corporation, Irvine, CA), and, as such, there is little detail in the literature regarding the signal processing aspects of its implementation. In contrast, the DPOP measure is well known, the author has developed a sophisticated algorithm for its measurement [30, 31] and

![Fig. 5](image1.png)

**Fig. 5** Relationship between DPOP and PMod. Linear resistors of 5/20/50 cmH2O/l are indicated by solid/dashed/dotted lines with circles/triangles/squares respectively. High/low breathing efforts are indicated by large/small symbols at the end of each line. All data were collected with a coached breathing at a respiration rate of 8 brpm.

![Fig. 6](image2.png)

**Fig. 6** Respiration effort induced change in DPOP and PPV using the modified DPOP algorithm. a Relationship between DPOP and PPV, b DPOP change versus PPV change.
has access to the proprietary hardware necessary to extract the raw signal required for its computation.

This work has shown that there is an increase in DPOP associated with an increasing effort to breathe. However, this increase is initiated at the subject’s current DPOP value which is subject-specific. How the DPOP value may be used to provide an absolute measure of respiratory effort is something for future work. However, it may have use as a trend indicator, i.e. as an indicator of improving or deteriorating patient respiratory status.

5 Conclusions

In the study reported here healthy volunteers were tasked to breathe over a range of respiratory efforts at fixed RR. The subject fluid levels were assumed constant over the short time periods involved and thus would not influence the DPOP value. The results indicate that the increased effort to breathe, in the form of an increase in mouth pressures while breathing against a resistance, manifests itself as an increase in DPOP. Further, a relationship between DPOP and PMod was found. In this regard a version of the DPOP algorithm that corrects for low perfusion was implemented which resulted in an improved relationship between DPOP and PPV.

Although a limited cohort of seven volunteers was used, the results suggest that DPOP may be useful as a respiratory effort parameter, given that the fluid level of the patient is maintained at a constant level over the period of analysis. No apparent difference in results were obtained between those obtained for constriction on inhalation and those for constriction on exhalation. However, this should be tested further with a larger cohort. A suggested next step in the examination of the DPOP parameter as a measure of respiratory effort would be to calculate it continuously during a therapeutic intervention for a respiratory issue: for example, a decrease in DPOP over time may be expected for an asthmatic patient given drug therapy to relieve symptoms. Although normally thought of as a fluid responsiveness parameter, the work suggests that DPOP may have a wider role in patient monitoring. As such, it may more properly be described in the terms first given to it by Cannesson et al. [21], i.e. as a measure of “respiratory variations in pulse oximetry plethysmographic waveform amplitude”. Depending on its mode of use, it may perform the function as a measure of fluid responsiveness, or as suggested by the current work, as a measure of respiratory effort.

Compliance with ethical standards

Conflicts of interest Paul S. Addison is an employee of Medtronic who sponsored the research.

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