Motion resistant pulse oximetry in neonates

R Sahni, A Gupta, K Ohira-Kist, T S Rosen

Background: Pulse oximetry is widely used in neonates. However, its reliability is often affected by motion artefact. Clinicians confronted with questionable oxygen saturation (SpO₂) values often estimate the reliability by correlating heart rate (HR) obtained with the oximeter with that obtained by electrocardiogram.

Objective: To compare the effects of motion on SpO₂ and HR measurements made with Masimo signal extraction technology and those made with a Nellcor N-200.

Design: Continuous pulse oximetry and HR monitoring were performed in 1.5 healthy, term infants (mean (SD) birth weight 3408 (458) g) undergoing circumcision, using Masimo and Nellcor pulse oximeters and a standard HR monitor (Hewlett-Packard). Simultaneous minute by minute behavioural activity codes were also assigned. Baseline data were collected for 10 minutes when the infant was quietly asleep and then continued during and after circumcision for a total duration of one hour. The oximeter HR and SpO₂ values were compared and related to HR values obtained by ECG during all three periods. The effect of behavioural activity on SpO₂ and HR was also evaluated.

Results: When compared with results obtained with the Nellcor, the mean SpO₂ and HR were higher and the incidence of artefact lower with the Masimo during all three periods. Masimo HR more accurately predicted HR obtained with a standard monitor, with lower residual error. SpO₂ and HR values obtained with the Nellcor were lower and more variable during all behavioural states, especially crying, when excessive motion artefact was most likely.

Conclusions: The data suggest that Masimo signal extraction technology may offer improvement in pulse oximetry performance, particularly in clinical situations in which extreme motion artefacts are likely.

METHODS

Study population

The study population consisted of 15 healthy, term male infants (mean (SD) birth weight 3408 (458) g), all of whom were participants in a randomised controlled study of impact of non-invasive analgesia on the response of the infant to circumcision. All studies were approved by institutional review board, and written consent was obtained from parents of all infants.

Experimental design

Continuous pulse oximetry and HR monitoring were performed using Masimo signal extraction technology and Nellcor N-200 pulse oximeters and a standard HR monitor (Hewlett-Packard 3680). Baseline data were collected for 10 minutes with the infant quietly asleep. Data collection was then continued during and after circumcision for a total duration of one hour. Simultaneous minute by minute assessments of behavioural sleep and activity state were also made.

Abbreviations: HR, heart rate; ECG, electrocardiogram; SpO₂, oxygen saturation
Experimental method
All circumcisions were performed in the infant physiology laboratory, located adjacent to the nursery. Infants were brought to the laboratory approximately half an hour before the procedure, at which time ECG leads were attached, and disposable adhesive pulse oximeter sensors (Nellcor N-25 and Masimo low noise optical probe Neo) were placed on the left foot. These sensors were optically shielded from each other to eliminate sensor to sensor optical cross talk. The probes were then connected to the corresponding oximeter. Baseline data were collected for 10 minutes when the infant was quietly asleep with minimal movements. Infants were then given the appropriate analgesia, restrained in the Circumstraint (Olympic Medical, Seattle, Washington, USA), and circumcision was performed. Data collection continued during and after the procedure for the total duration of one hour. Sleep state was coded during the entire study period. Behaviour codes were assigned each minute using a scoring system developed and validated in our laboratory. Active sleep was coded whenever at least one rapid eye movement was observed during the minute, and quiet sleep was designated when the infant was asleep without rapid eye movements. During quiet sleep the infant was relaxed and appeared “rag-doll” floppy with movements limited to startles and jaw jerks. An indeterminate state was coded when small body movements were observed, but no rapid eye movements were seen. Codes were also assigned for wakefulness and crying. 16 movements were observed, but no rapid eye movements were seen. The accuracy and reliability of conventional pulse oximetry is limited by subject motion. A possible explanation for this limitation involves the confounding effects of venous blood flow. Conventional pulse oximeters assume that the arterial compartment is the only compartment that contains pulsating blood flow. During patient motion, however, the venous efflux is also pulsing, giving the instrument a target that is not completely deoxygenated. This may cause an increase in the pulsating component of venous blood flow, which is not detected by the pulse oximeter. The accuracy of the oximeter may be reduced, and the heart rate may be overestimated. Different state behaviors were assigned each minute using a scoring system. The accuracy and reliability of conventional pulse oximetry is limited by subject motion. A possible explanation for this limitation involves the confounding effects of venous blood flow. Conventional pulse oximeters assume that the arterial compartment is the only compartment that contains pulsating blood flow. During patient motion, however, the venous efflux is also pulsing, giving the instrument a target that is not completely deoxygenated. This may cause an increase in the pulsating component of venous blood flow, which is not detected by the pulse oximeter. The accuracy of the oximeter may be reduced, and the heart rate may be overestimated. Different state behaviors were assigned each minute using a scoring system. The accuracy and reliability of conventional pulse oximetry is limited by subject motion. A possible explanation for this limitation involves the confounding effects of venous blood flow. Conventional pulse oximeters assume that the arterial compartment is the only compartment that contains pulsating blood flow. During patient motion, however, the venous efflux is also pulsing, giving the instrument a target that is not completely deoxygenated. This may cause an increase in the pulsating component of venous blood flow, which is not detected by the pulse oximeter. The accuracy of the oximeter may be reduced, and the heart rate may be overestimated.
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Oxygen saturation (SpO₂) and heart rate (HR) recorded with Masimo and Nellcor pulse oximeters (artifact-free data) before, during, and after circumcision (n = 15)

Table 2

<table>
<thead>
<tr>
<th></th>
<th>Baseline</th>
<th>Circumcision</th>
<th>After circumcision</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Masimo</td>
<td>Nellcor</td>
<td>p Value</td>
</tr>
<tr>
<td>SpO₂ (%)</td>
<td>98.2 (0.3)</td>
<td>97.5 (0.3)</td>
<td>NS</td>
</tr>
<tr>
<td>HR (beats/min)</td>
<td>126.6 (3.0)</td>
<td>123.2 (3.6)</td>
<td>0.04</td>
</tr>
</tbody>
</table>

Values are mean (SD).

Blood also moves irregularly, which adds a confounding signal to the pulse waveform signal, and thereby changes the apparent amount of light transmitted to the photoreceptor at the two wavelengths, red and infrared. These are the two wavelengths that are used to detect the relative proportions of oxyhaemoglobin and deoxyhaemoglobin in the tissue (red/infrared transmitted light ratio). The magnitude of error in SpO₂ measurement will be influenced by venous blood saturation, arterial signal amplitude, and magnitude of motion. The motion-added signal produces false SpO₂ values to the extent that the extra signal, which does not represent the arterial pulse oxyhaemoglobin/deoxyhaemoglobin signal, is large relative to the arterial pulse signal. When the motion is major, absolutely or in relation to arterial pulse signal, the motion-added signal tends to predominate over the pulse signal, so that the red/infrared ratio transmitted to the photoreceptor produces a false SpO₂ value. An additional and specific source of error in conventional pulse oximetry occurs when noise from a variety of signal inputs that are not a part of the true arterial pulse signal is introduced equally into both the red and infrared channels, driving the red/infrared ratio towards unity. This ratio of 1.0 predicts a saturation of ~82% according to conventional pulse oximetry algorithm. Such motion and noise artefacts are disturbing. They also limit accurate assessment of oxygenation, particularly in the most ill patients, who are most susceptible to erroneous saturation readings.

Periods of absent or inaccurate SpO₂ values due to motion artefact are not insignificant, and may affect up to 50% of the recorded tracings in preterm and term infants. Motion artefact during pulse oximetry recordings is also dependent on behavioural state. Recent reports have documented as many as six changes in behavioural state per hour in preterm infants, with at least 10 changes that last more than 10 minutes per hour. Nursing interventions and noise independently result in significant changes in both the behavioural and physiological responses of infants studied. Such problems are worse during transport and during more physically active periods such as resuscitations. These are specific situations in which accurate pulse oximetry values would improve our understanding of abnormal oxygenation and the effects of oxygen therapy and other interventions.

Various methods have been devised to minimise the loss of signal associated with motion. Most of these involve averaging the signal over longer periods of time and/or rejecting rapid unphysiological changes in the signal ratio. Both techniques represent best guesses rather than true measurements. A new electronic signal processing technique and sensor design, called signal extraction technology by Masimo Corporation, has been reported to considerably decrease false SpO₂ values during patient motion. The Masimo paradigm is based on the assumption that arterial blood is not the only absorptive substance in tissue that may generate a pulsatile optical signal. An adaptive filter system is used to separately identify the energy present at each saturation in the range 1–100% in 0.5% increments every 0.4 second. Once the discrete saturation transformation identifies and verifies the energy peak produced by the pulsating arterial blood, which is usually the peak corresponding to the highest saturation in the discrete saturation transformation, it reports this value as the patient’s SpO₂. Thus, it is able to cancel the noise signal and separate the arterial signals from artefacts. This results in superior signal acquisition and reduces the risk of false alarms during motion at the monitoring site.

Our study shows that the SpO₂ and HR signals from Masimo are less variable, and the Masimo HR signal more accurately predicts the Hewlett-Packard HR during both resting and increased activity states. With signal extraction technology, the superior signal handling during increased activity states associated with higher degree of motion artefact is understandable, but differences are also observed during baseline and behavioural quiet sleep periods. Various factors could potentially contribute to these differences. The signal drop out rate—that is, periods with no acceptable data—is much higher with conventional oximetry and may contribute most significantly to these differences. Of course, the internal smoothing algorithms and other editing functions limit the impact of these disturbances on the digital output of both devices, and the zero outputs are invisible to the observer. For example, in our study, during quiet sleep, which is associated with greatly reduced but not totally absent physical activity, the drop out rate in the raw data was 50 times greater for the Nellcor than the Masimo oximeter (4.16% vs 0.09%). An artefact reduction algorithm was applied to the raw data before the comparative analysis, and, despite that, the performance of the Masimo oximeter was noted to be superior. Another possible factor that may have contributed to these differences is sensor to sensor cross talk, as

Table 3

Relation of Masimo and Nellcor oximeter heart rates to cardiac monitor heart rate before, during, and after circumcision (n = 15)

<table>
<thead>
<tr>
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<th>Baseline</th>
<th>Circumcision</th>
<th>After circumcision</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Masimo</td>
<td>Nellcor</td>
<td>Masimo</td>
</tr>
<tr>
<td>Correlation coefficient (r²)</td>
<td>0.86</td>
<td>0.63</td>
<td>0.72</td>
</tr>
<tr>
<td>Residual error (beats/min)</td>
<td>1.3</td>
<td>3.9</td>
<td>5.8</td>
</tr>
<tr>
<td></td>
<td>0.78</td>
<td>0.31</td>
<td>2.4</td>
</tr>
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both oximeter sensors were on the same foot. During the planning of this experimental protocol, we had noted that the leg movements were not symmetrical when the infants were restrained in the Circumstraint, particularly during times of excessive motion (circumcision). This movement asymmetry would have significantly affected our primary objective of comparing the oximeter performance during motion. So, to test the oximeter performance under a similar degree of motion, the sensors were placed on the dorsum of the left foot as far from each other as possible and optically shielded to prevent cross talk. However, this may not have completely eliminated it, and cross talk may have contributed to minor differences in $SpO_2$; this is a study limitation.

Our data suggest that the Masimo signal extraction technology may improve pulse oximetry performance particularly in clinical situations in which extreme motion is likely, such as circumcision and crying in our neonates. Also, superior performance during times of minimal motion or when the infant is motionless suggests better signal acquisition and processing with the Masimo signal extraction technology. Further studies investigating the effects of this new technology on clinical care and outcomes will be necessary. Such application studies should include evaluation of nursing time and effort, medical judgments about routine oxygen deficits and oxygen therapy, and assessments of usefulness during active behaviour, transport, and resuscitation. Instrumentation that is considerably more accurate and reliable, with much greater sensitivity and specificity, may offer a definite advantage over current home monitoring systems, which have notoriously high rates of false alarms. A more accurate and reliable pulse oximeter should have the capacity to refine the therapeutic use of oxygen and benefit patient management and outcome.

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**REFERENCES**