ORIGINAL RESEARCH

FREQUENCY AND DURATION OF OXIMETER DROP-OUTS IN THE NICU: AN OBSERVATIONAL STUDY

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Abstract

Oximeters used for continuous monitoring experience periods with no signal. This SpO2 drop-out is widely acknowledged and its causes generally understood. This is a prospectively designed analysis of an existing database with the aim of characterizing drop-outs as experienced in the neonatal ICU. The data reflects 116 days of monitoring in seven tertiary care neonatal ICUs in 6 countries. From the evaluation of 1,396 drop-outs we found that typically the time was minimal with missing SpO2 data, and the episodes were short (median 2.79 minutes per day IQR 0.17–76, median 22 seconds IQR: 15–37, respectively). During about half of the days there were no prolonged dropouts (1 minute or longer), even so half of the total time spent with no SpO2 data were in prolonged episodes (median length 110 seconds IQR 85–150). The predominate factor associated with excessive drop-out time was the number of prolonged episodes rather than their duration. We concluded that the impact of drop-outs during manual control of inspired oxygen primarily impact alarm fatigue, but that during automatic FiO2 control they could have an important impact. The relative effectiveness of the fall-back strategies of these automatic control systems ought to be evaluated.

Keywords

oximeter, oxygen saturation, neonatal ICU, SpO2 drop-out

Background

Continuous monitoring of neonatal oxygenation using the pulse oximeter is an essential tool in the neonatal intensive care unit (NICU). Relatively small changes in overall exposure to high and low extremes of peripheral oxygen saturation (SpO2) levels have been shown to have significant impact on neonatal outcome [1, 2]. In addition, exposure to extreme levels have been shown to be primarily episodic with episodes of 1 minute or longer contributing most significantly to bad outcomes [3].

While an essential tool, the use of the pulse oximeter is problematic. False alarms are common and known to create significant alarm fatigue [4]. It is well documented that this leads to risks of frequent disregard for alarm policy and SpO2 targeting guidelines [5].

The reasons for these false readings, that is artifact, are understood. They are primarily caused by motion, low regional perfusion and loss of sensor integrity. Even though the SpO2 is calculated based on an average of 3–15 pulses, there are still occasions when no reading is presentable. In this occasion it is said that the SpO2 reading “drops out”. While this is a common occurrence, we are unaware of any reports characterizing drop-outs in the NICU.

Much of the practice of adjusting the inspired oxygen using pulse oximetry is applied pragmatically by the attending nurse. General good practice would dictate that oxygen adjustments should be made judiciously, with a goal of avoiding responses to false positives or providing unnecessary extra oxygen in response to a short transient physiological perturbation. Periods of drop-outs are treated in the same manner. However, new neonatal ventilators offering nearly continuous automated adjustment of FiO2 based on monitored SpO2 have been shown to be quite effective [6]. International standards do require fail-back modes for periods of missing or unreliable signals. “Fail-back” being the preferred term to the commonly used, “fail-safe”.

As a part of our work to improve alarm and oxygenation control practices, we are evaluating a large database of SpO2 recordings from NICUs. As part of this EPI-SAT project, we supposed it important to characterize the frequency and duration of SpO2 drop-outs as they occur in real NICUs.
**Methods**

The EPI-SAT database includes 116 days of continuous SpO$_2$ data from 58 significantly preterm infants receiving respiratory support and supplemental oxygen. These infants all participated in studies of SpO$_2$ control and are from seven neonatal ICU’s in six countries [7, 8]. Consent was received for participation in the studies and all the data in the EPI-SAT database is de-identified.

SpO$_2$ readings were collected every 5 seconds, by a data-logger attached to the digital output of a mechanical ventilator (AVEA-CLiO2, Vyaire Medical, Irvine, CA, USA). The oximeter used Masimo SET technology (Masimo, Irvine, CA, USA) with averaging set at 8 seconds. Purpose built software (MatLab, MathWorks, MA, USA) evaluated drop-out episodes and parsed them into 5 duration categories: < 5 s, 5 to < 30 s, 30 to < 60 s, 60 to < 120 s, 120 to < 600 s. Episodes of 10 minutes or longer were excluded as likely procedural interventions, rather than drop-outs. The latter two categories (length of 1 minute or greater) were defined as being relevant to SpO$_2$ control (FiO$_2$ < 30 s, 21% for 30 to < 60 s, 21% for 60 to < 120 s, and 21% for 120 to < 600 s). The longest drop-outs. When present, the median duration of these prolonged drop-outs was almost 2 minutes (110 s, IQR 85–150).

**Results**

We identified and analyzed 1,396 drop-outs during the 116 recorded days, that is 12 per day. The typical total time with missing SpO$_2$ readings (drop-outs) was minimal (median 2.79 minutes/day, range 0.17–76), obviously skewed. The actual median duration of individual drop-outs was 22 seconds (IQR: 15–37).

The distribution of drop-out time among the five duration categories was: 7% for < 5 s, 24% for 5 to < 30 s, 21% for 30 to < 60 s, 21% for 60 to < 120 s, and 29% for 120 s or greater. Thus about half of the drop-out time was deemed, according to our prospective definition, relevant to SpO$_2$ control. Details of the variation among the duration categories are tabulated in Table 1. During 59 of the 116 days, there were no prolonged drop-outs. When present, the median duration of these prolonged drop-outs was almost 2 minutes (110 s, IQR 85–150).

**Excessive drop-out time (outliers with >8.6 minutes of drop-out in a day) occurred in only 14 of the 116 days. In this group of outliers the median duration of time without SpO$_2$ readings was 16 minutes (IQR 11–31), significantly longer than the nominal group (2 minutes (IQR 1–4), p<0.001). The predominant factor in the difference between the two categories was the number of prolonged episodes, rather than their duration. The median duration of prolonged drop-out episodes in the outlier drop-out group was 134 s (IQR 8–147) and was not significantly different from nominal group: 100 s (IQR 80–145). In contrast, the median rates of prolonged drop-outs per day were 4 (IQR 3.25–9.25) and 0 (IQR 0–1) respectively, and the difference was significant (p < 0.001). Other details of the difference in the frequency of prolonged drop-outs are tabulated by outlier category in Table 2.**

**Discussion**

In this group of very preterm infants who required respiratory support and supplemental oxygen, oximeter dropouts were common, that is one every few hours. Furthermore, the total time without SpO$_2$ readings was relatively low and the typical duration of each episode relatively short. Only about half the days saw one or more prolonged drop-out, and only 12% of the days experienced an atypically high amount of time with

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**Table 1: Total Drop-Out time in each duration category (minutes/day).**

<table>
<thead>
<tr>
<th>Category</th>
<th>n (n/day)</th>
<th>Median (IQR)</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 5 s</td>
<td>485 (4.2)</td>
<td>0.17 (0.08–0.25)</td>
<td>0–18</td>
</tr>
<tr>
<td>5 to &lt; 30 s</td>
<td>572 (4.9)</td>
<td>0.83 (0.42–1.33)</td>
<td>0–29</td>
</tr>
<tr>
<td>30 to &lt; 60 s</td>
<td>195 (1.7)</td>
<td>0.58 (0–1.35)</td>
<td>0–15</td>
</tr>
<tr>
<td>60 to &lt; 120 s</td>
<td>89 (0.8)</td>
<td>0 (0–1.35)</td>
<td>0–16</td>
</tr>
<tr>
<td>120 to &lt; 600 s</td>
<td>55 (0.5)</td>
<td>0 (0–2.0)</td>
<td>0.2–32</td>
</tr>
</tbody>
</table>

**Table 2: Frequency of Episodes 1 minute or longer (n/24 hours).**

<table>
<thead>
<tr>
<th>Category</th>
<th>n</th>
<th>Median (IQR)</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>typical</td>
<td>102</td>
<td>0 (0–1)</td>
<td>0–3</td>
</tr>
<tr>
<td>outliers</td>
<td>14</td>
<td>4 (3.25–9.25)</td>
<td>2–14</td>
</tr>
</tbody>
</table>

The differences are statistically significant (p < 0.001).
Auto-FiO2, when compared to manual control, reduces the risk of excessive overshoot in oxygen concentration. Predictably, it appears that following drop-outs and the results are promising [personal communication].

One such study has just finished of these Auto-FiO2 controllers is not documented. One impact and must be addressed. In general the approach of commercially available systems ought to be documented. Additional studies to broaden these findings and these results are warranted.

Conclusion

We confirmed that oximeter drop-outs probably do not generally impact clinical decision making during manual titration of inspired oxygen. While their prevalence is limited, and they are of short duration, they do contribute to false alarms and the potential for alarm fatigue as an acknowledged risk factor. In contrast, we suggest their prevalence would have an impact on the effectiveness during automated control of inspired oxygen. For this reason the fall-back approaches of commercially available systems ought to be documented and evaluated.

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Author contributions

The first author (TB) provided the concept for the study, and wrote the first draft of the manuscript. The second author (KR) coded the data extraction software. Both authors critically reviewed the data and the manuscript.

References


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