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Case Study

Influence of the duration of noninvasive ventilation on the cardiorespiratory indicators of preterm infants: A randomized clinical trial

Abstract

This study aimed to evaluate the cardiorespiratory indicators of preterm infants submitted to two noninvasive ventilator support systems in two periods. It was an controlled randomized clinical trial (RBR-79579d). Fifty-two newborns (gestational age of 30.6±2.3 weeks, weight of 1,366±445 g) submitted to continuous positive airway pressure (CPAP) therapy were studied. The infants were randomly allocated to two groups: 31 children received CPAP therapy for 48 hours and 21 children for 72 hours. The respiratory rate, heart rate, oxygen saturation and Silverman-Anderson Score were recorded once a day. Three measurements were obtained in a 15 minutes intervals. The infants were monitored during the predetermined period of noninvasive ventilation and for 24 hours after pressure support withdrawal. The Student t-test was used for comparison of the variables between groups after normality evaluation by the Kolmogorov-Smirnov test and significance was considered when p<0.05. It was found that the birth characteristics were homogeneous, without significant differences between groups. The cardiorespiratory indicators did not differ significantly between groups, but better mean values were observed for the 72-hour group. There was a difference in the respiratory rate of 5.0 breaths per minute and the heart rate was 7.8 bpm, while oxygen saturation was similar in the two groups.

Conclusions: Neither the duration of noninvasive ventilation of 48 or 72 hours nor the nasal CPAP and nasal intermittent positive pressure had an influence on the cardiorespiratory indicators of preterm infants. However, the period of 72 hours resulted in lower respiratory and heart rates.

Abbreviations

CPAP: Nasal Continuous Positive Airway Pressure; SAS: Silverman-Anderson Score

Introduction

Nasal continuous positive airway pressure (CPAP) has been valued as a beneficial strategy in neonatal intensive care units. Since nasal CPAP is a noninvasive ventilation method, it reduces the inflammatory response in the lung parenchyma [1], decreases the failure rates of extubation [2] and reintubations [3], minimizes lung injury [4-6] and reduces respiratory problems such as apnea and respiratory acidosis, as well as the use of supplemental oxygen [7]. Despite the known benefits of nasal CPAP in preterm infants, there is a lack of specific knowledge about the parameters adopted [8], modalities used [9] and effects of different interfaces [10,11]. The duration of noninvasive ventilation is also a factor of interest since the literature has shown important variability in the effects of different durations [12,13].

The duration of nasal CPAP in newborns, a lower gestational age and birthweight are directly related to the risk of nasal trauma [14,15]. So far, no studies have addressed the relationship between the duration of noninvasive ventilation and cardiorespiratory response. Part of the scientific studies have focused on the relationship the ventilator support and body position [16-18].

Out hypothesis was that cardiorespiratory indicators in premature newborns had better values when premature infants are submitted to longer periods of noninvasive support and nasal intermittent positive-pressure in better mode.

The purpose of the present study was to evaluate the cardiorespiratory indicators of preterm infants submitted to two noninvasive ventilator support systems in two periods.

Patients And Methods

This was a single-blind, two-arm parallel group, randomized clinical trial conducted at the Infant Intensive Care
Center of a public tertiary hospital. The study was approved by the Ethics Committee of the same institution (No. 070/2009) and was registered in the Brazilian Clinical Trials Registry (RBR-79d9th). The infants were included in the study after the parents or legal guardians had signed the free informed consent form.

Eighty preterm infants with a birthweight ranging from 540 to 2,450 g and a gestational age of 24 to 36 weeks, who received noninvasive ventilation support (CPAP) through a silicon nasal prong (Hudson® RCI Infant Nasal Prong CPAP cannula system, Teleflex, Inc., USA), were eligible. Children with heart diseases, congenital malformations, hydrocephalus and surgical indications, and newborns who did not complete the selected period of nasal CPAP were excluded.

**Study protocol**

Noninvasive ventilation was administered with a mechanical breathing system consisting of an expiratory valve for the control of positive expiratory pressure and a mixer for the oxygen fraction. For nasal CPAP, a continuous flow of 7 to 8 l/min, a positive expiratory pressure of 5 cmH2O, and an oxygen fraction of 21 to 40% were used. The following parameters were adopted for CPAP with nasal intermittent positive pressure: continuous flow (7 to 8 l/min), positive expiratory pressure (5 cmH2O), inspiratory pressure (15 cmH2O), controlled respiratory rate (14 breaths per minute), time of inspiration (0.6 s), and oxygen fraction (21 to 40%).

The children were allocated randomly to the groups using sealed envelopes containing the identification of the method (Nasal CPAP and Nasal intermittent positive-pressure) stratified into two times (48 and 72 hours).

The following cardiorespiratory variables were collected: respiratory rate determined by observing the number of chest movements over one minute timed with a chronometer, heart rate and oxygen saturation evaluated noninvasively with a Dixtal monitor (DX2010), and respiratory distress using the Silverman-Anderson Score (SAS), which ranges from 0 (no respiratory distress) to 10 (maximum respiratory distress). The parameters analyzed were: grunting, nasal flaring, intercostal retractions, xiphoid retractions, and synchrony of chest and abdominal movements.

The variables were collected once a day at the same time. Three measurements were obtained at intervals of 15 minutes. The infants were monitored during the predetermined period of ventilatory support (48 and 72 hours) and 24 hours after pressure support withdrawal.

This was a non–probability sample since the group had homogenous birth characteristics. The birth characteristics are reported descriptively as the mean, standard deviation, and relative frequency. The Student t-test was used for comparison of the variables between groups after normality evaluation by the Kolmogorov-Smirnov test and WilcoxonMann–Whitney test for non-parametric variables. All analyses were performed using the GraphPad InStat software (version 3.06 for Windows) and significance was considered when p<0.05.

### Results

During the study, 28 newborns were excluded because they did not complete the preestablished period of noninvasive ventilator support. The final sample consisted of 52 preterm infants that received noninvasive ventilatory support [Figure 1].

The two groups submitted to 48 and 72 hours of noninvasive ventilation were homogenous in terms of their birth characteristics, with no significant difference in the variables analyzed (Weight, gestational age, male, antenatal corticoid, previous mechanical ventilation and surfactant), p>0.5. The only variable that differed significantly between groups, but without a direct influence on the outcome, was the frequency of cesarean delivery, which was 21.3% higher in the 72-hour group (p=0.027).

In general, comparison during and in the first 24 hours after pressure support withdrawal showed no significant difference in the respiratory rate between the 48- and 72-hour groups or between modalities. However, mean respiratory rates tended to be lower in the 72-hour group. Separate analysis showed a significant difference in the respiratory rate between the 48-hour and the 72-hour cycling CPAP groups in the first 24 hours after pressure support withdrawal (p=0.029), with a difference of 5.2 breaths per minute in the 72-hour group. A significant difference during and after noninvasive ventilation was only observed when the 48-hour CPAP group was compared to the 72-hour cycling CPAP group. The difference between mean respiratory rates was about 6 breaths per minute, with a lower rate in the 72-hour cycling CPAP group [Table 1].

The maximum boletim Silverman Anderson did not exceed 1.5 and was similar in the groups monitored. A significant difference was only observed for the 72-hour CPAP group when the cycling and conventional groups were compared after weaning (p=0.024), but this difference had no clinical repercussions.

![Figure 1: Flow diagram of preterm newborns that were randomly allocated to receive nasal intermittent positive pressure and conventional nasal CPAP for 48 and 72 hours. The 28 newborns were excluded because they did not complete the predetermined period of noninvasive ventilation.](image)

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The heart rate showed a similar behavior when the periods of 48 and 72 hours and the nasal CPAP modalities (with and without cycling) were compared. The mean heart rates were within the expected range for age and did not result in undesired clinical repercussions [Table 2].

Oxygen saturation did not differ significantly between infants submitted for 48 and 72 hours to nasal CPAP and nasal intermittent positive pressure for 48 and 72 hours. There was also no difference when the period of noninvasive ventilation and the first 24 hours after weaning were compared. A significant difference was only observed between the groups submitted to CPAP for 48 hours with and without intermittent positive pressure (p=0.001). However, the mean oxygen saturation values were similar in the groups and within the normal range. All groups exhibited desirable mean oxygen saturation levels [Table 3].

**Discussion**

Noninvasive ventilation of newborns with respiratory distress before extubation is a recognized and valued practice in intensive care units. However, there are no reports in the scientific literature investigating the influence of the duration of noninvasive ventilation on neonatal cardiorespiratory indicators. Evidence indicates wide variation ranging from hours to weeks in the duration of nasal CPAP among newborns [12,13].

The relationship between a longer duration of the CPAP device in the nostrils and nasal trauma has been recognized [19]. The occurrence of nasal bleeding has been shown to be directly proportional to the duration of nasal CPAP therapy in preterm infants, with the observation of a higher risk after 48 hours of therapy [13].

**Table 1:** Mean respiratory rate of preterm infants submitted to nasal CPAP and nasal intermittent positive.

| CPAP 48 vs CPAP 72 h | Overall mean (breaths per minute) | % Difference between means |
|----------------------|----------------------------------|---------------------------|
|                      | % Difference 95% CI p            |
| During               | 50.6x48.5 2.1 (-)2.82-7.02 0.380 |
| After Int CPAP 48 vs Int. CPAP 72 h | 52.1x50.7 1.4 (-)7.4.10.20 0.766 |
| During               | 50.3x44.4 5.9 (-)1.55-13.35 0.118 |
| After                | 50.9x45.7 5.2 0.60-9.80 0.029 |
| CPAP 48 vs Int. CPAP 72 h | During 50.6x44.4 -6.0 (-)2.11-(-)9.88 0.003 |
|                      | After 52.1x45.7 -7.0 (-)3.38-(-)10.6 0.000 |
| CPAP 48 vs Int. CPAP 48 h | During 50.6x50.3 0.3 (-)6.79-7.39 0.923 |
|                      | After 52.1x50.9 1.2 (-)3.86-6.26 0.643 |
| CPAP 72 vs Int. CPAP 72 h | During 48.5x44.4 4.1 (-)2.81-11.01 0.233 |
|                      | After 50.7x45.7 5.0 (-)2.30-12.30 0.228 |

**Table 2:** Mean heart rate of preterm infants submitted to nasal CPAP and nasal intermittent positive pressure for 48 and 72 hours.

| CPAP 48 vs 72 h | Overall mean (bpm) | % Difference between means |
|----------------|--------------------|---------------------------|
|                | % Difference 95% CI p |
| During         | 144.7x136.9 7.8 (-)1.78-17.38 0.104 |
| After Int. CPAP 48 vs Int. CPAP 72 h | 142.7x149.4 (-)6.7 (-)20.7-7.31 0.325 |
| During         | 142.1x141.0 1.1 (-)8.46-10.6 0.818 |
| CPAP 48 vs Int. CPAP 72 h | During 144.7x141.0 (-)3.0 3.53-(-)9.53 0.356 |
| After          | 142.7x143.9 1.0 (-)11.46-13.46 0.871 |
| During         | 144.7x14 2.1 2.6 (-)6.96-12.1 0.584 |
| After          | 149.4x143.9 5.5 (-)12.21-23.1 0.520 |

CPAP continuous positive airway pressure, Int CPAP intermittent positive pressure, BPM beats per minute, 95% CI 95% confidence interval, H hours.

**Table 3:** Mean oxygen saturation of preterm infants submitted to nasal CPAP and nasal intermittent positive pressure for 48 and 72 hours.

| CPAP 48 vs 72 h | Overall mean (%) | % Difference between means |
|----------------|-----------------|---------------------------|
|                | Difference 95% CI p |
| During         | 96.5x95.9 0.6 (-)0.89-2.09 0.434 |
| After Int. CPAP 48 vs Int. CPAP 72 h | 96.5x97.0 (-)0.5 (-)1.89-0.89 0.470 |
| During         | 94.6x96.8 (-)1.5 (-)5.59-2.59 0.450 |
| After CPAP 48 vs Int. CPAP 72 h | 97.7x96.8 0.9 0.04-1.6 0.077 |
| During         | 96.5x96.1 0.0 (-)0.72-0.72 1.00 |
| After CPAP 48 vs Int. CPAP 48 h | 96.5x96.8 0.0 (-)0.72-0.72 1.00 |
| During         | 96.5x94.6 1.9 (-)2.36-6.16 0.363 |
| After CPAP 72 vs Int. CPAP 72 h | 96.5x97.7 (-)1.2 (-)1.87-(-)0.53 0.001 |
| During         | 95.9x96.1 (-)0.2 (-)1.60-1.20 0.772 |
| After          | 97.0x96.8 0.2 (-)1.35-1.75 0.878 |

CPAP continuous positive airway pressure, Int CPAP intermittent positive pressure, 95% CI 95% confidence interval, H hours.

In the present study investigating the duration and modality of noninvasive ventilation, we found no important difference in the mean values of the cardiorespiratory variables studied. Thus, the infants showed a similar behavior during the two periods and in the two modalities.

Although the cardiorespiratory variables were within the desired range for age in the present study, infants submitted to noninvasive ventilation for 72 hours tended to have lower mean respiratory rates, with a difference of up to 5.9 breaths per minute during noninvasive ventilation. The same trend was observed in the first 24 hours after weaning from the pressure support, with a difference of up to 5.2 breaths per...
minute. There was also a trend towards lower respiratory rates in the 72-hour intermittent positive pressure CPAP group. The sample studied exhibited mean SAS scores less than 1.5 and the lowest mean score was observed in the 72-hour intermittent positive pressure CPAP group (0.6), demonstrating that the infants were not experiencing respiratory distress. This result can be explained by the influence of positive expiratory pressure through noninvasive ventilation. However, the infants exhibited a low Boletim Silverman Andersen even after pressure support withdrawal, irrespective of the duration of noninvasive ventilation.

Respiratory rate responses of preterm infants have been demonstrated under different clinical conditions. Studies have shown lower mean respiratory rates in newborns in the prone position before and after feeding [20], during tube feeding [21], in oxygen-dependent infants [22], and during quiet sleep [23]. In the present study, despite the lack of significant differences between groups, lower respiratory rates were observed during the period of 72 hours of noninvasive ventilation and in the intermittent positive pressure modality.

The literature indicates that the heart rate also responds to neonatal conditions. Lower mean heart rates of newborns have been demonstrated in the supine position, during active sleep [24], during spontaneous breathing [25] and during ventilator support [17]. However, the infants studied here exhibited mean heart rates expected for age. The noninvasive ventilation modality and the duration of pressure support did not influence the heart rate of the groups. The threshold of 170 bpm was observed in only one infant during noninvasive ventilation and in five infants after withdrawal of the pressure support. The latter may have been due to support withdrawal and not to the modality and/or duration of ventilation.

Oxygen saturation did also not differ between the groups submitted to noninvasive ventilation for 48 and 72 hours. The oxygen mixer was part of the nasal CPAP system and the inspiratory fraction used ranged from 21 to 40%. This inhaled fraction of oxygen may have directly influenced the oxygen saturation values obtained, since there was no effect of the duration of noninvasive ventilatory support. Another important factor is the positive expiratory pressure provided by the nasal CPAP prong, which increases the alveolar area and consequently the functional residual capacity, optimizing gas exchange and increasing saturation [13]. It was therefore expected that oxygen saturation was influenced by these factors. However, the infants continued to exhibit desirable values after weaning from noninvasive ventilation and no significant difference was observed between groups.

However, some limitations of the study should be mentioned. The sample size of each separate group was small and the results therefore only apply to this profile of preterm infants. Furthermore, the infants received pressure support for 48 and 72 hours, a period during which the respiratory system achieves greater balance of ventilatory mechanics. Future studies should therefore include a group submitted to noninvasive ventilation for 24 hours.

Conclusion

In conclusion, the results suggest no advantages of noninvasive ventilation for 48 and 72 hours or of the intermittent positive pressure and conventional modality of nasal CPAP in preterm infants. However, the period of 72 hours resulted in lower respiratory and heart rates.

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Authors’contributions

Carmen Silveira, participated in the development of the protocol, contributed to the writing of the manuscript, had primary responsibility for protocol development, patient screening, enrollment, outcome assessment and writing the manuscript Kamila Leonardi and Ana Paula Melo, participated in the development of the protocol, had primary responsibility for protocol development, patient screening, enrollment, outcome assessment José Zaia, analytical framework for the study Marisa Brunherotti, supervised the design and execution of the study, performed the final data analyses and contributed to the writing of the manuscript.

All authors read and approved the final manuscript

Ethics approval

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

The study was approved by Ethics Committee of the University of Franca (number 070/2009, Feb 2010) and was registered in the Brazilian Clinical Trials Registry (RBR-7d9dh).

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