Noninvasive Ventilation Strategies for Early Treatment of RDS in Preterm Infants: An RCT

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BACKGROUND AND OBJECTIVES: There is evidence that new methods of noninvasive ventilation (NIV) support have significantly changed respiratory distress syndrome (RDS) management in preterm infants. Further perspectives for neonatologists involve the assessment of different NIV strategies in terms of availability, effectiveness, and failure. This study evaluates the efficacy of 2 different NIV strategies for RDS treatment in very low birth weight (VLBW) infants: nasal synchronized intermittent positive pressure ventilation (NSIPPV), which is a modality of conventional ventilation with intermittent peak inspiratory pressure, and bilevel continuous positive airway pressure (BiPAP), not synchronized, with 2 alternate levels of continuous positive airway pressure.

METHODS: We conducted a 2-center randomized control study in 124 VLBW infants (<1500 g and <32 weeks of gestational age) with RDS who received NIV support (NSIPPV, n = 62; BiPAP, n = 62) within 2 hours of birth. We evaluated the performance of NIV strategies by selected primary outcomes (failure rate and duration of ventilation) and secondary outcomes.

RESULTS: The number of failures and duration of ventilation support did not differ between NSIPPV and BiPAP strategies (P > .05 for both). Moreover, no differences between groups were found regarding secondary outcomes (P > .05 for all).

CONCLUSIONS: The present data show no statistically significant differences between NSIPPV and BiPAP strategies in terms of duration of ventilation and failures, suggesting that both NIV techniques are effective in the early treatment of RDS in VLBW infants. Further randomized investigations on wider populations are needed to evaluate the effect of NIV techniques on long-term outcomes.

WHAT’S KNOWN ON THIS SUBJECT: Noninvasive ventilation (NIV) reduced the need of intubation in preterm infants with RDS. However, randomized studies comparing nasal synchronized intermittent positive pressure ventilation and bilevel continuous positive airway pressure are still lacking.

WHAT THIS STUDY ADDS: The present study shows no differences in short-term outcomes between 2 different NIV strategies, nasal synchronized intermittent positive pressure ventilation and bilevel continuous positive airway pressure, in preterm infants for the initial treatment of RDS.
In recent decades, considerable changes have been made in the management of respiratory distress syndrome (RDS), supporting the notion that appropriate perinatal management can be effective by minimizing the use of mechanical ventilation (MV) in very low birth weight (VLBW) infants. In particular, antenatal steroid prophylaxis, accurate delivery room and respiratory management with early nasal continuous positive airway pressure (NCPAP), surfactant replacement in the early phase of RDS, the INSURE (intubation, surfactant extubation) procedure, and the increased use of noninvasive ventilation (NIV) have been shown to improve respiratory outcome.\textsuperscript{1–5}

The hypothetical advantages of NIV, compared with invasive MV, consist in the possibility to reduce barotrauma, biotrauma, and ventilator-induced lung injury. Data on NIV support, such as nasal intermittent positive pressure ventilation (NIPPV), nasal synchronized intermittent positive pressure ventilation (NSIPPV), and bilevel continuous positive airway pressure (BiPAP), are still controversial. On the one hand, NSIPPV/NIPPV has shown promising short- and long-term respiratory outcomes compared with NCPAP or MV.\textsuperscript{6–10} On the other hand, Kirpalani et al found no significant differences between NCPAP or NIV strategies (ie, NSIPPV/NIPPV/BiPAP) in a wider study population, in terms of mortality or occurrence of bronchopulmonary dysplasia (BPD).\textsuperscript{11} In this regard, Roberts et al described several discrepancies among studies previously conducted (ie, recruited populations, ventilation modalities, devices used, synchronization systems, and clinical applications) and concluded that, at this stage, no clear advantages were detectable for NIPPV or BiPAP over NCPAP in reducing mortality or BPD.\textsuperscript{12} Moreover, no studies elucidating any differences between NSIPPV and BiPAP, used as the primary mode of ventilation for RDS, are yet available, except for a nonrandomized study.\textsuperscript{13}

Therefore, the present randomized study aimed to investigate the effectiveness of these 2 different NIV strategies: NSIPPV, synchronized with an intermittent positive pressure, and BiPAP, nonsynchronized with 2 alternate levels of continuous positive airway pressure (CPAP), as the primary mode of ventilation in the respiratory management of RDS in VLBW infants in terms of duration and failure of NIV support and of selected secondary outcomes.

**METHODS**

We conducted a randomized study in 124 VLBW infants, admitted in 2 NICUs (C. Arrigo, Children’s Hospital, Alessandria, Italy, and V. Buzzi, Children’s Hospital, Milan, Italy) from January 2010 to December 2012, delivered before 32 weeks of gestational age (wGA) with a birth weight $\leq 1500$ g (Fig 1). Approval was obtained from the respective local ethics committees. Informed and written consent was obtained, before delivery, from all parents of the patients before inclusion in the study.

The protocol for delivery room management, RDS treatment, devices and interfaces used, and ventilator adjustment were the same for the 2 centers. Infants who had signs of RDS at birth were treated with sustained lung inflation (SLI)\textsuperscript{14} and NCPAP in addition to the American Academy of Pediatrics recommendations.\textsuperscript{15}

The respiratory strategy for RDS treatment in the newborns was as follows. In the delivery room, after oropharyngeal and nasal suctioning, pressure-controlled SLI (25 cmH$^2$O) was performed for 15 seconds using a neonatal mask and a T-piece ventilator (Neopuff Infant T-Piece Resuscitator, Fisher & Paykel, Auckland, New Zealand), followed by the delivery of 5 cmH$^2$O NCPAP.\textsuperscript{16} The SLI maneuver was repeated in patients in whom respiratory and/or heart failure persisted. After failure of the second SLI maneuver, infants were intubated. In both groups, neonatal care was started at the lowest oxygen concentration, between 0.21 and 0.4 fraction of inspired oxygen (FIO$_2$), to maintain arterial oxygen saturation (SaO$_2$) of 85% to 93%. All enrolled infants were transferred to the NICU with NCPAP support (5 cmH$^2$O). Further support depended on gestational age (GA):

**FIGURE 1**
Flow chart describing recruitment.
1. Newborns \( \leq 26 \) wGA received a prophylactic replacement of surfactant with INSURE in the first 2 hours of life. After INSURE, the infants were switched to either BiPAP or NSIPPV.

2. Newborns \( > 26 \) and \( \leq 29 \) wGA received BiPAP or NSIPPV in the first 2 hours of life.

3. Newborns \( > 29 \) wGA not requiring or positively responding to the initial resuscitation maneuver were maintained on NCPAP support until arrival at the NICU; after 2 hours from birth, if they did not show any signs of RDS, NCPAP was stopped. BiPAP/NSIPPV was performed if clinical and blood gas analysis patterns were suggestive of RDS as follows: (a) need of \( F_{O_2} > 0.4 \) and/or (b) \( pH < 7.20 \) and/or \( P_{O_2} < 50 \) mm Hg and/or \( P_{CO_2} > 65 \) mm Hg and (c) clinical patterns of RDS characterized by retractions and/or dyspnea. Apnea, defined as \( \geq 4 \) episodes/hour or need for mask ventilation \( \geq 2 \) times/hour, was another criterion to start NIV support.

Newborns complicated by RDS requiring NIV support within 2 hours from birth but not intubated were allocated by use of computer-generated random numbers to receive either NSIPPV \((n = 62)\) or BiPAP \((n = 62)\). Apneic or severely depressed newborns requiring MV within 2 hours from birth were excluded from the study and started on MV.\(^{17}\)

**BiPAP**

BiPAP support was delivered using the Infant Flow-driver device (Infant Flow System, ViaSys Corp, Yorba Linda, CA). We used the short binasal prongs as interface (CareFusion, Yorba Linda, CA) with different sizes according to weight. This method of nonsynchronized ventilation support provides 2 alternate levels, lower and higher, of CPAP; the newborn can breathe spontaneously on these 2 levels to create 2 different functional residual capacities (FRCs). The initial ventilator parameters were lower and higher, CPAP levels 4 to 6 cmH\(_2\)O and 8 to 9 cmH\(_2\)O, respectively; a \( t_{n_{\text{eigh}}} \) of 1 second; and a pressure exchange rate of 20/minute, with the lowest adjusted \( F_{O_2} \) to maintain an \( S_{O_2} \) of 88% to 93%. Respiratory settings (CPAP lower maximum 7 cmH\(_2\)O, CPAP higher maximum 10 cmH\(_2\)O, pressure exchange rate max 30/minute) were adjusted to guarantee blood gas analysis within normal ranges. Weaning was started with a progressive reduction of the set pressure exchange rate (minimum 15 pressure exchanges/minute), followed by a reduction of the higher CPAP level to 6 cmH\(_2\)O and the lower CPAP level to 4 cmH\(_2\)O. BiPAP was stopped when infants showed no signs of RDS with \( F_{O_2} < 0.30 \).

**NSIPPV**

NSIPPV support was delivered with a nasal ventilator device (Giulia, Ginevri, Rome, Italy) that detects the inspiratory effort by means of a pneumotachograph equipped with a fixed orifice (2-mm diameter for low birth weight infants). This is a conventional strategy of synchronized ventilation provided by nasal interface, with short binasal prongs (NIV set, Ginevri, Rome, Italy) of different sizes according to weight. The inspiratory flow was detected as a pressure change across the resistance, positioned proximally to the nasal interface. The initial ventilator parameters were positive-end expiratory pressure (PEEP) 4 to 6 cmH\(_2\)O; peak inspiratory pressure (PIP) 15 to 20 cmH\(_2\)O; inspiratory time 0.3 to 0.4 second; flow rate 6 to 10 L/minute; respiratory rate (RR) 40 breaths per minute with the lowest adjusted \( F_{O_2} \) to maintain an \( S_{O_2} \) of 88% to 93%. Respiratory settings (PIP maximum 25 cmH\(_2\)O, PEEP maximum 7 cmH\(_2\)O, RR maximum 60 breaths per minute) were adjusted to guarantee blood gas analysis within normal ranges. The highest trigger sensitivity avoiding autotriggering was selected. Weaning from NSIPPV was performed with a reduction of RR to 15 breaths per minute with a PIP of 10 to 15 cmH\(_2\)O and a PEEP of 4 cmH\(_2\)O and was stopped when infants showed no signs of RDS with \( F_{O_2} < 0.30 \).

**Failure Criteria**

NIV failure was defined when 1 or more of the following criteria persisted or recurred, after a surfactant dose (maximum 3 doses) or within 12 hours from previous administration: (1) hypoxia (\( F_{O_2} \) requirement >0.40), (2) acidosis (\( pH < 7.20 \) and hypercarbia (\( P_{CO_2} > 65 \) mm Hg), and (3) apnea as \( \geq 4 \) episodes/hour or the need for mask ventilation \( \geq 2 \) times/hour. We also considered necrotizing enterocolitis (NEC), bowel perforation, and hemodynamic instability conditions for NIV failure.\(^{18}\)

**Surfactant Administration**

According to the protocol of respiratory management, the first dose of surfactant was administered at 200 mg/kg (Curosurf, Chiesi, Parma, Italy). Additional doses of surfactant were given at 100 mg/kg, at least 12 hours after previous administration. After INSURE, newborns received the same NIV support device as before.

All newborns were treated with caffeine (caffeine citrate 20 mg/kg loading dose; 5 to 10 mg/kg/day maintenance).

**Monitoring Parameters**

Newborns were monitored by using pre-postductal \( S_{O_2} \) monitoring (Masimo Datascipe Radical, Masimo Corporation, Irvine, CA). For each infant, the following variables were recorded: GA, BW, gender, main maternal pregnancy diseases, mode of delivery, and Apgar scores at 1 to 5 minutes. At study entry, \( F_{O_2} \), mean airway pressure (MAP), \( S_{O_2} \), pH, and \( P_{CO_2} \) were recorded.

**Primary Outcomes**

The primary end points were the duration of NIV support and failure rate.
Secondary Outcomes

Secondary end points were duration of respiratory support, incidence of pneumothorax (PNX), occurrence of moderate/severe BPD, incidence of intraventricular hemorrhage (IVH) more than second degree, periventricular leukomalacia (PVL), need for second/multiple surfactant doses, need for postnatal glucocorticoid treatment, persistence of patent ductus arteriosus (PDA) requiring pharmacological treatment, retinopathy of prematurity (ROP) more than second degree, NEC, late-onset sepsis (LOS), death, and days to regain BW. Moderate/severe BPD was defined according to the classification of Jobe and Bancalari,19 IVH was classified according to Papile et al,20 and ROP was graded according to the criteria established by the International Committee for Classification of ROP.21

Statistical Analysis

For the calculation of sample size, we used duration of ventilation as the main primary outcome. As no basic data are available for this high-risk population, we were able to retrieve the duration of ventilation by the 2 different NIV strategies from the database of our 2 NICUs. These data were used for the power calculation. We assumed a difference of 24 hours between the 2 groups in the duration of NIV as clinically relevant. At a confidence level \( \alpha = 0.05 \) and power level of 0.80, we needed 62 patients for each group.22 The sample size was calculated by using nQuery Advisor (Statistical Solutions, Saugus, MA), version 5.0.

Data were reported as means and SD and median and interquartile ranges for continuous variables, whereas absolute and relative frequencies were used for categorical variables. Parameters of the 2 groups were compared using Student \( t \) or Mann-Whitney \( U \) 2-sided tests for continuous variables and \( \chi^2 \) or Fisher exact test for categorical variables.

\( P < .05 \) was considered statistically significant, and all \( P \) values were based on 2-tailed tests. Statistical analysis was performed by using SPSS for Windows (SPSS, Chicago, IL).

RESULTS

Table 1 shows the perinatal characteristics in the studied groups. No significant differences (\( P > .05 \)) were found between NSIPPV and BiPAP groups for wGA, BW, gender, incidence of cesarean delivery, premature rupture of membranes, evidence of chorioamnionitis, occurrence of pregnancy hypertension requiring antihypertensive agent treatment, abruptio placenta, occurrence of multiple pregnancies, complete course of prenatal glucocorticoids prophylaxis, and Apgar scores at 1 and 5 min. No differences were shown regarding blood gas analysis, \( \text{FiO}_2 \), \( \text{SaO}_2 \), pH, and \( \text{Pco}_2 \) at study entry. We observed a significant difference in MAP that was higher in the NSIPPV group than the BiPAP group, depending on the initial ventilator settings.

Table 2 shows primary and secondary outcome characteristics. No significant differences were found between groups in terms of duration of ventilation on NIV support and incidence of failure. Moreover, there were no significant differences in the incidence of postnatal death, moderate/severe BPD, PNX, IVH, PVL, postnatal glucocorticoid administration, multiple surfactant doses, PDA, ROP, NEC, LOS, or time to regain BW. PNX occurred in 6 cases (NSIPPV, \( n = 2 \); BiPAP, \( n = 4 \)) and represented a cause of NIV failure in 3 cases (NSIPPV, \( n = 2 \); BiPAP, \( n = 1 \)).

In 18 of 124 infants (NSIPPV, \( n = 10 \); BiPAP, \( n = 8 \)), NIV support failed. The causes were early-onset sepsis (NSIPPV, \( n = 5 \); BiPAP, \( n = 3 \)), pulmonary hypertension (NSIPPV, \( n = 5 \); BiPAP, \( n = 4 \)), hypoxia, and hypercapnia (NSIPPV, \( n = 2 \); BiPAP, \( n = 4 \)), hypoxia alone (NSIPPV, \( n = 3 \); BiPAP, \( n = 3 \)), PDA (NSIPPV, \( n = 1 \)), NEC (BiPAP, \( n = 2 \)), and PNX (NSIPPV, \( n = 2 \); BiPAP, \( n = 1 \)). The timing of NIV failure did not differ between studied groups (median [25th to 75th centile] for NSIPPV, 36 hours [17 to 72]; for BiPAP, 34 hours [17 to 72]).

### Table 1 Perinatal Characteristics of Preterm Infants Supported by NSIPPV or BiPAP

<table>
<thead>
<tr>
<th></th>
<th>NSIPPV (n = 62)</th>
<th>BiPAP (n = 62)</th>
<th>( P )</th>
</tr>
</thead>
<tbody>
<tr>
<td>BW, g</td>
<td>1108 ± 276</td>
<td>1165 ± 275</td>
<td>.23</td>
</tr>
<tr>
<td>GA, wks</td>
<td>28.6 ± 2.1</td>
<td>28.8 ± 2.2</td>
<td>.66</td>
</tr>
<tr>
<td>GA ≤26 wks</td>
<td>9</td>
<td>9</td>
<td>.999</td>
</tr>
<tr>
<td>GA &gt;26 and ≤29 wks</td>
<td>33</td>
<td>32</td>
<td>.999</td>
</tr>
<tr>
<td>GA &gt;29 wks</td>
<td>20</td>
<td>21</td>
<td>.999</td>
</tr>
<tr>
<td>Male/female</td>
<td>27/35</td>
<td>25/37</td>
<td>.85</td>
</tr>
<tr>
<td>Small for GA</td>
<td>14</td>
<td>12</td>
<td>.82</td>
</tr>
<tr>
<td>Cesarean delivery</td>
<td>38</td>
<td>47</td>
<td>.12</td>
</tr>
<tr>
<td>Preterm premature rupture of membrane</td>
<td>13</td>
<td>20</td>
<td>.22</td>
</tr>
<tr>
<td>Chorioamnionitis</td>
<td>10</td>
<td>12</td>
<td>.63</td>
</tr>
<tr>
<td>Pregnancy-induced hypertension requiring treatment with antihypertensive agents</td>
<td>13</td>
<td>20</td>
<td>.22</td>
</tr>
<tr>
<td>Abruptio placenta</td>
<td>10</td>
<td>10</td>
<td>.999</td>
</tr>
<tr>
<td>Twins</td>
<td>10</td>
<td>15</td>
<td>.37</td>
</tr>
<tr>
<td>Prenatal steroids completed course</td>
<td>55</td>
<td>57</td>
<td>.76</td>
</tr>
<tr>
<td>Apgar score at 1 min</td>
<td>7 ± 1</td>
<td>7 ± 1</td>
<td>.999</td>
</tr>
<tr>
<td>Apgar score at 5 min</td>
<td>8 ± 1</td>
<td>8 ± 1</td>
<td>.999</td>
</tr>
<tr>
<td>( \text{SaO}_2 ) at study entry, %</td>
<td>86 ± 12</td>
<td>87 ± 14</td>
<td>.67</td>
</tr>
<tr>
<td>( \text{FiO}_2 ) requirement at study entry</td>
<td>0.32 ± 0.04</td>
<td>0.35 ± 0.05</td>
<td>.78</td>
</tr>
<tr>
<td>MAP at study entry, mmHg</td>
<td>7.6 ± 1.0</td>
<td>6.2 ± 1.2</td>
<td>&lt;.01</td>
</tr>
<tr>
<td>pH at study entry</td>
<td>7.21 ± 0.15</td>
<td>7.23 ± 0.10</td>
<td>.38</td>
</tr>
<tr>
<td>( \text{Pco}_2 ) at study entry, mmHg</td>
<td>54 ± 9</td>
<td>53 ± 8</td>
<td>.51</td>
</tr>
</tbody>
</table>

Data are presented as the mean ± SD or n.
TABLE 2 Primary and Secondary Outcomes in Preterm Infants Supported by NSIPPV or BiPAP

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>NSIPPV (n = 62)</th>
<th>BiPAP (n = 62)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Primary outcomes</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nasal ventilation support, h</td>
<td>89 (61–145)</td>
<td>87 (48–134)</td>
<td>.45</td>
</tr>
<tr>
<td>Failure of nasal ventilation support</td>
<td>10</td>
<td>8</td>
<td>.80</td>
</tr>
<tr>
<td><strong>Secondary outcomes</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Death</td>
<td>0</td>
<td>2</td>
<td>.49</td>
</tr>
<tr>
<td>Moderate/severe BPD</td>
<td>7</td>
<td>7 (n = 60)</td>
<td>1.00</td>
</tr>
<tr>
<td>PNX</td>
<td>2</td>
<td>4 (n = 60)</td>
<td>.43</td>
</tr>
<tr>
<td>IVH &gt;2nd degree</td>
<td>2</td>
<td>2 (n = 60)</td>
<td>1.00</td>
</tr>
<tr>
<td>PVL</td>
<td>2</td>
<td>4 (n = 60)</td>
<td>.43</td>
</tr>
<tr>
<td>Postnatal glucocorticoids</td>
<td>5</td>
<td>5 (n = 60)</td>
<td>1.00</td>
</tr>
<tr>
<td>PDA</td>
<td>18</td>
<td>14 (n = 60)</td>
<td>.54</td>
</tr>
<tr>
<td>ROP &gt;2nd degree</td>
<td>2</td>
<td>3 (n = 60)</td>
<td>.68</td>
</tr>
<tr>
<td>NEC</td>
<td>0</td>
<td>2 (n = 60)</td>
<td>.49</td>
</tr>
<tr>
<td>Multiple surfactant doses</td>
<td>21</td>
<td>18 (n = 60)</td>
<td>.97</td>
</tr>
<tr>
<td>Surfactant, &gt;26 wGA</td>
<td>19 (n = 53)</td>
<td>21 (n = 52)</td>
<td>.78</td>
</tr>
<tr>
<td>Early-onset sepsis</td>
<td>13</td>
<td>15 (n = 60)</td>
<td>.67</td>
</tr>
<tr>
<td>LOS</td>
<td>21</td>
<td>14 (n = 60)</td>
<td>.23</td>
</tr>
<tr>
<td>Time to regain BW, d</td>
<td>14 ± 4</td>
<td>13 ± 4</td>
<td>.17</td>
</tr>
</tbody>
</table>

Data are presented as the median (25th to 75th centile), n, or mean ± SD.

**DISCUSSION**

In the last decade, new therapeutic strategies and technological advances have considerably changed RDS treatment in VLBW infants. New delivery room management and early NIV support significantly contributed to a sensible decay in the need for MV support.23–28 Thus, further perspectives for neonatologists involve the assessment of different NIV strategies in terms of availability, effectiveness, and failure.

In the present 2-center randomized study, we found no differences in primary and secondary end points between 2 different NIV strategies (ie, NSIPPV and BiPAP) performed as primary modes for RDS treatment. Results are consistent and offer additional support to a previous nonrandomized observation using NSIPPV and BiPAP as primary modes in the treatment of RDS.13 In our series, we also found a low incidence of failure (18 of 124 newborns, 15%) and a brief time of respiratory support (median for NSIPPV 89 hours; for BiPAP 87 hours). Moreover, no correlations were found between failure occurrences and GA subgroups.

Low failure in NIV support can be also explained on the basis of perinatal treatments, such as prenatal glucocorticoid prophylaxis (85% to 90% for our population), known to be effective on lung immaturity, and improvements in delivery room management such as SLI and early NCPAP. Recent observations reported an improved postnatal adaptation, in terms of lung and cardiovascular function, in SLI-treated infants and animals.29,30 Another explanation can be the early NCPAP support in the delivery room, which is known to be beneficial for lung outcome.2,23–28

Although the current study shows that both methods of NIV (SIPPV and BiPAP) are feasible and probably equally effective, it does not answer the question whether NIV is better than NCPAP as primary treatment of RDS. Several authors in smaller study populations (NIPPV or BiPAP versus NCPAP) and a meta-analysis reported less need of MV, less risk of intubation in the first 72 hours from birth, and reduction of hospitalization duration and O2 dependency.7,10,31–33

However, Kirpalani et al, in a recent large multicenter trial, showed no significant differences in terms of mortality or BPD occurrence between NCPAP and NIV strategies, used both as first intention or in the weaning phase, but without a specific protocol for NIV (devices, modalities, synchronization).11 Finally, Roberts et al suggested that NIPPV (synchronized or nonsynchronized) might be advantageous over NCPAP as primary support for reduction of hours [19 to 65]) (P > .05) (Table 3).

We did not find statistically significant differences in the incidence of failure either between the 2 study groups (P > .05 for all) or between failure subgroups after stratification for wGA (Table 3). In addition, infants who failed did not significantly differ in baseline characteristics from infants who did not fail on NIV.

TABLE 3 Characteristics, Timing, and Causes of Failure of Preterm Infants Who Failed on NIV

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>NSIPPV (n = 10)</th>
<th>BiPAP (n = 8)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>BW, g</td>
<td>1000 ± 510</td>
<td>980 ± 268</td>
<td>.92</td>
</tr>
<tr>
<td>GA, wks</td>
<td>28 ± 1</td>
<td>28 ± 1</td>
<td>.93</td>
</tr>
<tr>
<td>GA ≤26 wks/total GA subgroup</td>
<td>2/9</td>
<td>2/9</td>
<td>1.00</td>
</tr>
<tr>
<td>GA &gt;26 and ≤28 wks/total GA subgroup</td>
<td>7/33</td>
<td>4/32</td>
<td>.34</td>
</tr>
<tr>
<td>GA &gt;29 wks/total GA subgroup</td>
<td>1/20</td>
<td>2/21</td>
<td>.96</td>
</tr>
<tr>
<td>Apgar score at 1 min</td>
<td>6 ± 1</td>
<td>6 ± 1</td>
<td>1.00</td>
</tr>
<tr>
<td>Apgar score at 5 min</td>
<td>8 ± 1</td>
<td>8 ± 1</td>
<td>1.00</td>
</tr>
<tr>
<td>Prenatal steroids completed course</td>
<td>7</td>
<td>6</td>
<td>.77</td>
</tr>
<tr>
<td>Timing of failure (median, 25°–75° centile), h</td>
<td>36 (17–72)</td>
<td>34 (19–65)</td>
<td>.85</td>
</tr>
<tr>
<td>Early-onset sepsis</td>
<td>5</td>
<td>3</td>
<td>.96</td>
</tr>
<tr>
<td>Hypoxemia</td>
<td>3</td>
<td>3</td>
<td>.87</td>
</tr>
<tr>
<td>Hypcapnia and hypoxemia</td>
<td>2</td>
<td>4</td>
<td>.32</td>
</tr>
<tr>
<td>Persistent pulmonary hypertension of the newborn</td>
<td>5</td>
<td>4</td>
<td>1.00</td>
</tr>
<tr>
<td>PNX</td>
<td>2</td>
<td>1</td>
<td>1.00</td>
</tr>
<tr>
<td>PDA</td>
<td>1</td>
<td>0</td>
<td>1.00</td>
</tr>
<tr>
<td>NEC</td>
<td>0</td>
<td>2</td>
<td>.47</td>
</tr>
</tbody>
</table>

Data are presented as the mean ± SD, n, or median (25th to 75th centile).
intubation, although there is no clear advantage.12

On the basis of the present findings, bearing in mind that NIV strategies can act through different modalities, further investigations evaluating their effectiveness in RDS treatment are justified.34 Indeed, NSIPPV uses a conventional synchronized modality of intermittent positive pressure ventilation, delivered through a nasal interface, whereas BiPAP uses a nonsynchronized ventilation that provides alternately 2 different CPAP levels without intermittent peak of inspiratory pressure, in which newborns can breathe spontaneously, creating 2 different FRCs. Both methods have theoretical benefits. In particular, (1) NSIPPV through intermittent increase in pressure enhances tidal volume (Vt), minute ventilation, and MAP, resulting in better alveolar recruitment and gas exchange9,12,35; (2) BiPAP, by using a much longer timehigh permits a complete respiratory cycle (inspiration and expiration) on the higher CPAP level, creating 2 different FRCs; increases MAP; and FRC switching generates a Vt with better gas exchange.7,36 Additional common NIV advantages, due to pressure changes, consist in the stimulation of spontaneous breathing that reduces failure risk due to apneas.37 Another issue deserving further consideration concerns NIV settings. On the one hand, several authors highlight the need for strict protocols and guidelines6–38; on the other hand, there is still no consensus, since conflicting results have been reported in terms of variability of airway pressure from the set pressure, increase in Vt and minute ventilation, and the drop in work of breathing.39–43 Therefore, further studies are needed to evaluate the transmission of setting pressures to the lower airways in an open system with large and variable leakage and improve NIV synchronization systems with infant spontaneous breathing. However, we found no difference in effectiveness between nasal-flow synchronized (NSIPPV) and nonsynchronized (BiPAP) strategies.

Data on prophylactic/early surfactant administration available at the time of patient recruitment were still controversial and under debate. Therefore, in infants ≤26 wGA, we chose prophylactic surfactant administration (ie, within 2 hours from birth, after stabilization on NCPAP) for the higher risk of failure,23 whereas a selective therapeutic strategy was planned for those >26 wGA.44 Currently, prophylactic approaches do not seem to be justified, and further investigations to clarify the efficacy of early NIV with the best timing for surfactant administration, especially in extremely low birth weight infants, are eagerly awaited.45 We did not find any differences between groups in the need for surfactant single/multiple doses.

In the present series, we found no statistically significant differences in secondary outcomes between the 2 NIV devices. Of course, the small number of infants eligible for statistical analysis of secondary outcomes does not allow us to draw definite conclusions. In this respect, we observed a moderate/severe BPD incidence, comparable to that of previous studies.38,46 The low incidence of PNX, NEC, or bowel perforations suggests that NIV techniques could be considered reasonably safe for these infants.

Last but not least, successful NIV management requires a high quality of neonatal care. High-risk infants require experienced nurses for the best cleaning of the upper airways, nasal cannula positioning, and maintaining the containment position of the newborns. These precautions are implemented to ensure effective airflow, maintaining adequate pressure from the nostrils to the distal airways, to increase comfort of the newborns and prevent trauma to the nostrils.47

CONCLUSIONS
The present data show that both NSIPPV and BiPAP, used as primary respiratory support in the treatment of RDS of VLBW infants, are feasible and equally effective. These results prompt further RCT investigations to evaluate the effectiveness of different NIV strategies on long-term outcomes.

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