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Spontaneous breathing and respiratory support of preterm infants at birth

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CHAPTER 7

Early respiratory management of respiratory distress syndrome in very preterm infants and bronchopulmonary dysplasia: a case-control study

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Abstract

Background: Immediately after birth, preterm infants are highly susceptible to lung injury. Early nasal continuous positive airway pressure (CPAP) is an attempt to avoid intubation and may minimize lung injury. In contrast, early nasal CPAP can fail and at that time surfactant rescue can be less effective.

Objective: To compare the pulmonary clinical course and outcome of very preterm infants (gestational age 25-32 weeks) with respiratory distress syndrome (RDS) who started with early nasal CPAP and failed (ECF group), with a control group of infants matched for gestational age, who were directly intubated in the delivery room (DRI group). Primary outcome consisted of death during admission or bronchopulmonary dysplasia (BPD)

Results: 25 infants were included in the ECF group and 50 control infants matched for gestational age were included in the DRI group. Mean gestational age and birth weight in the ECF group were 29.7 weeks and 1,393 g and in the DRI group 29.1 weeks and 1,261 g ($P = ns$). The incidence of BPD was significantly lower in the ECF group than in the DRI group (4% vs 35%; $P < 0.004$; OR 12.6 (95% CI 1.6-101)). Neonatal mortality was similar in both groups (4%). The incidence of neonatal morbidities such as severe cerebral injury, patent ductus arteriosus, necrotizing enterocolitis and retinopathy of prematurity, was not significantly different between the two groups.

Conclusion: A trial of early nasal CPAP at birth may reduce the incidence of BPD and does not seem to be detrimental in very preterm infants. Randomized controlled trials are needed to test whether early respiratory management of preterm infants with RDS plays an important role in the development of BPD.

Introduction

Early respiratory management specifies ventilatory support from birth until stabilization during the first days of life, starting with resuscitation in the delivery room. However, the recently updated international guidelines on neonatal resuscitation do not provide specific recommendations on how to provide respiratory support in very preterm infants due to lack of data (1). Various strategies are currently used to compensate for inadequate respiratory drive, surfactant deficiency and/or disturbances in the reabsorption of lung liquid after birth. Early nasal continuous positive airway pressure (CPAP), started directly after birth, is a gentle and effective non-invasive attempt to support non-compliant lungs (2-6). It has been shown in animal studies that early nasal CPAP minimizes lung injury after preterm birth (7;8). During the immediate postnatal period the preterm, surfactant-deficient lung is highly susceptible to tissue injury and an exaggerated, unchecked inflammatory reaction can be triggered with the first manual breaths during resuscitation (9-13). Several cohort studies report less requirement of mechanical ventilation when early nasal CPAP is used (14). Nonetheless, a considerable percentage of preterm infants with respiratory distress syndrome (RDS) fails early nasal CPAP and requires intubation and mechanical ventilation during the first days of life (15;16). During this time period a potential improvement of lung function by surfactant treatment may have been missed as surfactant-deficient preterm infants respond better to surfactant treatment when it is given early in the course of their disease (17;18).

The aim of this case-control study was to compare the pulmonary course and outcome of very preterm infants with RDS who failed early nasal CPAP and a control group of infants, matched for gestational age, who were intubated in the delivery room.

Patients and Methods

Patients

All inborn very preterm infants with a gestational age ≤ 32 weeks admitted to the neonatal intensive care unit of the Leiden University Medical Center from January 2003 until December 2005 were identified. Limits of viability in the Netherlands are set at 25 weeks' gestation. Clinical data were collected from the hospital records. All infants with RDS who were supported with early nasal CPAP but failed and were intubated and ventilated within 72 hours after birth (Early CPAP Failure = ECF group) were selected for the study group. Each infant in the ECF group was matched for gestational age (± 1 week of gestation) with the next two consecutively born infants with RDS who were intubated in the delivery room (Delivery Room Intubation = DRI group). Diagnosis of RDS was based on clinical symptoms and chest X-ray findings. The following perinatal data were collected: birth weight, gestational age, gender, race, singleton or twin, maternal pregnancy disease, preterm premature rupture of the membranes, signs of chorioamnionitis (defined as maternal fever with at least

one of the following symptoms: leucocytosis, tenderness of the uterus, fetal tachycardia, foul-smelling amniotic fluid), mode of delivery, intra-uterine growth retardation, antenatal steroids (defined as any administration of steroids irrespective of completion of therapy), umbilical arterial pH, resuscitation in the DR (none, oxygen administration, positive pressure ventilation with mask and bag, intubation and mechanical ventilation), 5 min Apgar score. Postnatal data collection included: early nasal CPAP (defined as nasal CPAP support within 15 minutes after admission to the neonatal intensive care unit), intubation within 72 hours, time and cause of intubation, clinical signs of RDS and staging by chest X-ray, duration of mechanical ventilation (first period and total period), type of ventilation (conventional ventilation only, rescue high frequency oscillation), surfactant treatment, administration of postnatal steroids, and occurrence of air leaks.

Primary outcome consisted of death during admission or bronchopulmonary dysplasia (BPD) based on the NICHD definition (19). Secondary outcome was based on the presence of cerebral pathology detected by cranial ultrasound, retinopathy of prematurity, patent ductus arteriosus, and necrotizing enterocolitis.

Respiratory management

If respiration was insufficient in the delivery room, positive-pressure ventilation was started with a self-inflating bag and mask during 30 seconds. At first inflation, pressures till 30-40 cmH₂O could be given, thereafter not more than 20 cmH₂O was allowed. If signs of respiratory distress persisted, the decision to intubate and start mechanical ventilation or to start early nasal CPAP (within 15 minutes) was dependant on RDS severity. After arrival in the neonatal intensive care unit, Infants on nasal CPAP were placed on an Infant Flow Device (EME Tricomed, Brighton, United Kingdom) or Infant Star ventilator (Infrasonics Inc, San Diego, CA) with a nasopharyngeal tube. CPAP-levels were titrated from 5 to 8 cmH₂O according to the degree of respiratory distress, assessed by observing the infant's chest retraction, effort of breathing, chest radiograph appearance, and oxygen requirement (stabilize oxygenation until arterial PO₂ >50 mmHg, while pH >7.20 and arterial PCO₂ <60 mmHg). Caffeine or theophylline treatment was started as soon as possible after birth in neonates <30 weeks of gestation and only after apnea emerged in more mature infants.

Intubation and mechanical ventilation were initiated when infants exhibited arterial hemoglobin saturations <88% or arterial PO₂ < 50 mmHg while receiving FiO₂ ≥ 0.40 (corresponding with an AaPO₂ ratio <0.22) and/or arterial PCO₂ >60 mmHg, pH <7.20, or when more than 4 apnea episodes occurred in one hour or mask ventilation was needed >2 times per hour. Surfactant was given when on mechanical ventilation with a mean airway pressure x FiO₂ ratio >2. Infants with RDS were extubated as soon as a FiO₂ <0.3 and a mean airway pressure <7 cmH₂O was reached. Immediately after extubation nasal CPAP was started. Nasal CPAP was discontinued when the neonate remained stable with a capillary PCO₂

<60 mmHg, oxygen saturation >92% without supplementary O₂ and no CPAP for more than four consecutive hours. No changes in respiratory management occurred in the study period.

Data analysis

Data are presented as mean + SD or number (percentage) and were analyzed with SPSS software (SPSS for windows, version 12.0, 2005, Chicago, IL). Analyses compared infants who failed early nasal CPAP treatment and were intubated within 72 hours of age (ECF group) and matched infants who were intubated immediately in the DR (DRI group). The null hypothesis was that the ECF group did not have a better pulmonary clinical course and outcome than the DRI group. The baseline characteristics of the two groups were compared using Student's t test for parametric and the Mann-Whitney U test for non-parametric comparisons for continuous variables, and the χ^2 test for categorical variables. Reported p-values are two-sided; the level of significance is <0.05.

Results

Early respiratory management

A total of 268 inborn very preterm infants (gestational age range between 25 and 32 weeks) were admitted consecutively to the neonatal intensive care unit between January 1, 2003 and December 31, 2005. Six infants were excluded because of severe congenital anomalies of the cardiac or respiratory system or anomalies incompatible with survival. Mean birth weight was 1318 ± 385 g, mean gestational age 29.3 ± 1.9 weeks, and overall mortality 7% (19/268). The incidence of moderate or severe BPD among survivors was 15% (37/243). Early respiratory management of the 262 included infants is shown in Figure 1. Among the 208 infants who needed respiratory support, 42% received early nasal CPAP and 58% were intubated. Early nasal CPAP was started after arrival at the NICU in 83/87 (95%) infants and 4 (5%) of them received early nasal CPAP immediately in the DR. A nasopharyngeal tube was used in 85% of the cases and an infant flow driver in 15%. There were no significant differences in clinical failure between the two devices. The distribution of DRI, early nasal CPAP and ECF failure, plotted against gestational age is shown in Figure 2. Failure of CPAP occurred in 1/9 (11%) infants < 28 weeks, in 15/50 (30%) infants of 28-30 weeks and in 12/30 (40%) infants > 30 weeks of gestation. RDS was the principal cause of early nasal CPAP-failure (25/28 (89%) infants)

ECF vs. DRI in very preterm neonates with RDS

RDS was diagnosed in 127/262 (48%) very preterm infants, 83 of them (65%) were immediately intubated in the delivery room (DRI-group). In 42/127 (33%) infants with RDS early nasal CPAP was started within 15 minutes after birth, in 13 of them (31%) mask and bag

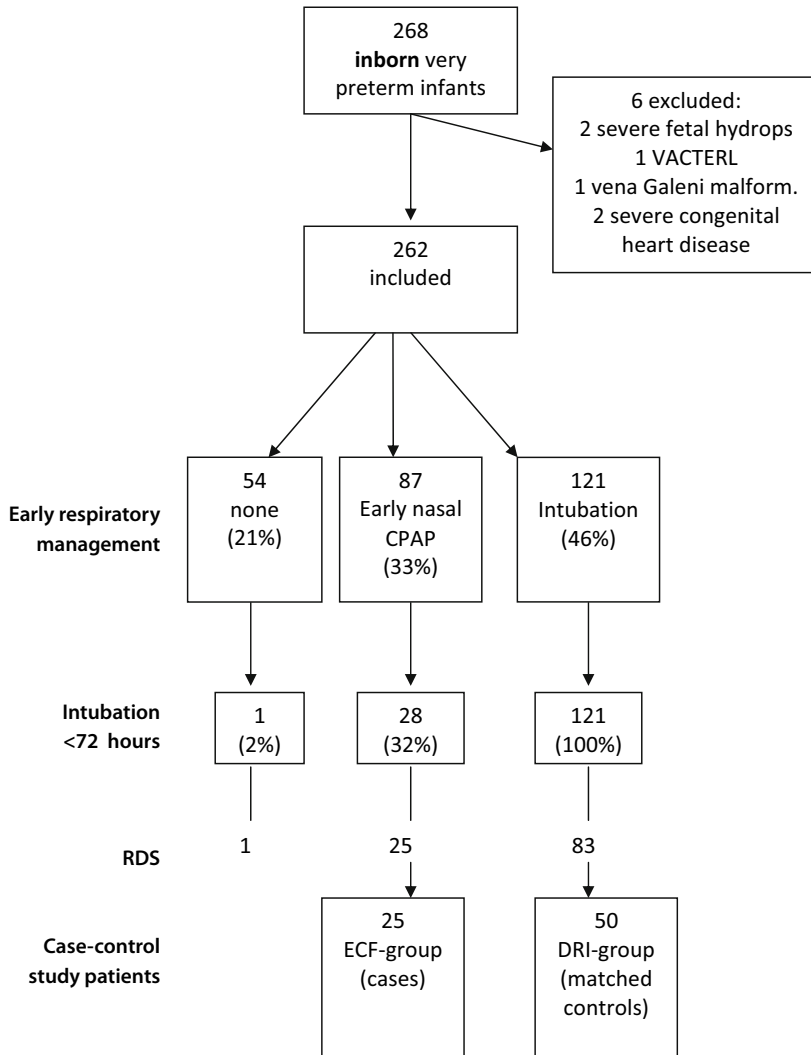


Figure 1. The distribution of need of ventilatory support in very preterm neonates (25-32 weeks of gestation). Early nasal CPAP= early nasal continuous positive airway pressure; mask and bag: using a self-inflating bag.

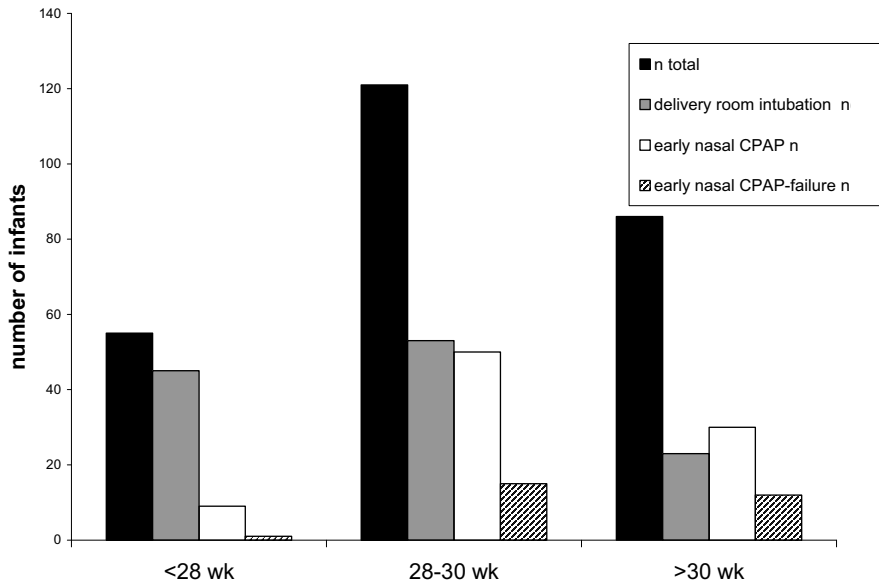


Figure 2. Distribution of delivery room intubation (DRI), early nasal continuous positive airway pressure and early nasal continuous positive airway pressure failure plotted against gestational age.

ventilation was given in the delivery room. In 25/42 (59%) infants early nasal CPAP failed within 72 hours of age and they needed endotracheal intubation, ventilation and surfactant treatment (ECF-group) (mean time of intubation 14 hours after birth; range 0.3-65 hours). The basic characteristics of the ECF-group and matched infants in the DRI-group were not significantly different (Table 1). Table 2 presents the pulmonary clinical parameters of the ECF and DRI groups. All patients with RDS intubated within 72 hours received surfactant, but significant more doses of surfactant were needed in the DRI-group.

Primary outcome is shown in Table 3. Survivors in the DRI group had significantly more often moderate to severe BPD at 36 weeks of gestation than survivors in the ECF group. Mortality was 4% in both groups and all died due to respiratory failure.

Comparison of secondary outcome between the ECF and DRI groups did not show significant differences in severe intraventricular hemorrhage (>grade 2) (3/24 (12.5 %) vs. 3/49 (6.1%)), cystic periventricular leucomalacia (0/24 (0%) vs. 2/47 (4%)) and patent ductus arteriosus requiring medical or surgical closure (5/25 (20%) vs. 15/50 (30%)). Necrotizing enterocolitis requiring surgery and severe retinopathy of prematurity was absent in both groups.

Table 1. Comparison of characteristics of infants with RDS who failed treatment with early nasal CPAP (ECF-group) and matched infants who were intubated immediately after birth in the delivery room (DRI-group).

Characteristic	Group ECF N = 25	Group DRI N = 50	Univariate analysis (P)
BW, g	1393 ± 379	1261 ± 395	NS
GA, wk	29.7 ± 1.6	29.1 ± 1.6	NS
Gender, male n (%)	9 (36)	27 (54)	NS
Umbilical arterial pH	7.28 ± 0.13	7.22 ± 0.13	NS
Prenatal steroids, n (%)	17 (68)	33 (66)	NS
Chorioamnionitis, n (%)	2 (8)	5 (10)	NS
PPROM, n (%)	6 (24)	15 (30)	NS
Toxaemia, n (%)	6 (24)	16 (32)	NS
Fetal distress, n (%)	9 (36)	18 (36)	NS
IUGR, n (%)	4 (16)	11 (22)	NS
Caesarean section, n (%)	12 (48)	24 (48)	NS
Apgar, 5 min < 7, n (%)	1 (4)	10 (20)	NS
Singletons, n (%)	13 (52)	20 (40)	NS

BW=birth weight; GA=gestational age; PPRM= preterm premature rupture of the membranes; IUGR= intra- uterine growth retardation.

Table 2. Pulmonary characteristics of neonates with RDS who failed early nasal CPAP treatment (ECF-group) and matched infants who were intubated in the delivery room (DRI-group).

Pulmonary clinical course parameters	ECF N = 25	DRI N = 50	Univariate analysis (P)
Surfactant, n doses	1.0 ± 0.5	1.6 ± 0.8	<0 .001
HFOV rescue, n (%)	3 (12)	15 (30)	NS
Total period of mechanical ventilation, days	3.9 ± 5.6	7.0 ± 9	NS
Pneumothorax, n (%)	1 (4)	3 (6)	NS
Dexamethason, n (%)	1 (4)	7 (15)	NS

CPAP=early nasal continuous positive pressure; RDS= respiratory distress syndrome; HFOV= high frequency oscillation ventilation.

Table 3. Primary outcome of infants who failed early nasal CPAP treatment (ECF-group) and matched infants who were intubated in the delivery room (DRI-group)..

Primary outcome	ECF N = 25	DRI N = 50	Univariate analysis (P)
BPD mild	7	11	NS
BPD moderate	1	14	0.01
BPD severe	0	3	NS
BPD _{mod-sev} , n (%)	1 (4)	17 (35)	0.004 OR 12.6 (95% CI 1.6-101)
Mortality, n (%)	1 (4)	2 (4)	NS

BPDmod-sev= moderate and severe bronchopulmonary dysplasia.

Discussion

Early nasal CPAP proved to be a valuable initial ventilatory strategy as a significant percentage of very preterm infants (33%) was successfully managed with early nasal CPAP alone, even extremely preterm infants (<28 weeks). However, early nasal CPAP can also fail, requiring infants to be intubated within 72 hours after birth, mostly because of RDS. This study showed that infants in the ECF group need less doses of surfactant and had a lower incidence of BPD compared to the matched DRI-infants. These findings are in agreement with animal studies, which have demonstrated that ventilatory management during the vulnerable period directly after birth may play an important role in the development of BPD (7;9-13;20). A trial of early nasal CPAP at birth may be a good alternative for intubation in case of RDS and may reduce the incidence of BPD. Because the ECF group does not have more adverse outcomes and complications than the DRI group, it seems reasonable to avoid or postpone intubation if possible, as long as early surfactant rescue is given if intubation becomes mandatory.

Initial and early respiratory management in preterm infants is a hot topic in neonatology. The IFDAS trial, presented and published in abstract form, gives a good representation of the main approaches (14). Many clinicians advocate elective intubation and early administration of surfactant (17;21). Others opt for an alternative approach of early nasal CPAP and selective surfactant treatment. Various non-randomized retrospective studies, with or without historic controls, report lower or no differences in rates of BPD in centers that try to avoid intubation and ventilation of preterm infants (21). However, the inclusion criteria, including gestational age of the preterm infants, the methods used to deliver nasal CPAP, and the levels of CPAP used vary between the studies. The interpretation of "early" nasal CPAP varies from starting in the delivery room directly after resuscitation to starting in the neonatal intensive care unit when signs of RDS appear (21). Early nasal CPAP and selective surfactant is an accepted alternative, but a Cochrane review of trials of prophylactic (= early) nasal CPAP recently concluded that there is currently insufficient information to evaluate the effectiveness of prophylactic nasal CPAP in very preterm infants (22). Retrospective studies in which early nasal CPAP is started in the delivery room have shown a reduction in incidence of BPD (3;4;6). To the best of our knowledge, this is the first case-control study comparing early nasal CPAP-failure and DR intubation in case of RDS. Similar to our study, Aly et al. retrospectively compared outcome of early nasal CPAP-failure and delivery room intubation in very preterm infants. However, they did not report the incidence of RDS in the total group and found, after controlling for basic characteristics, no difference in outcome, except for more necrotizing enterocolitis in the early nasal CPAP-failure group (16). We found no significant differences in secondary outcome, including necrotizing enterocolitis. Lindner et al. reported a significant decrease in delivery room intubation when a new ventilatory strategy was introduced, consisting of a continuous prolonged pressure-controlled inflation followed by CPAP at 4-6 cmH₂O (11). Their study was not designed to test the dif-

ference in outcome between direct intubation and early nasal CPAP-failure. Nevertheless, in case of RDS, the incidence of BPD was 40% when infants were intubated in the delivery room compared to 22 % when intubated later (6). In contrast to these reports and also to our findings, the infant flow driver and surfactant (IFDAS) trial found no reduction in BPD when early nasal CPAP was used. Lack of significance may have been the result of a lack of power due to randomisation of the 237 infants of 27-29 weeks of gestation to 4 treatment groups (14).

The period directly after birth is a very vulnerable phase and unnecessary intubation and ventilation may damage the upper airway and cause iatrogenic respiratory distress (13,20). The use of a self-inflating bag prior to intubation may also induce lung injury as positive end expiratory pressure can not be regulated and an accurate peak inspiratory pressure is difficult to achieve (23;24). Acute ventilator-induced lung injury sustained shortly after birth can trigger the inflammatory cascade, which may progress to BPD (7-13). Animals studies have shown that this inflammatory process may occur with the first manual breaths during resuscitation and compromise the effect of subsequent surfactant rescue treatment (9;10;12). Even administration of prophylactic surfactant does not protect against the adverse affects of large lung inflations at birth (11). In addition, lung injury after birth can be minimized by using early nasal CPAP instead of intubation and mechanical ventilation (7;8).

Symptoms of respiratory distress at birth in preterm infants may solely be due to their efforts to clear fluid from their lungs. Their ability to do so can be limited by surfactant deficiency and their soft and flexible chest wall makes preterm infants less able to generate sufficiently high pressures to achieve effective lung expansion (25;26). Nevertheless, expectant management (without direct intubation) may show that some preterm neonates are able to manage this with nasal CPAP assistance only. Once the alveoli are aerated, breathing needs less effort and distress symptoms may diminish. Some preterm infants who are intubated and ventilated in the DR may have had respiratory distress appropriate to a transitory phase, which eventually led to pulmonary complications due to "iatrogenic" RDS.

Compared to other reports, we found a higher percentage of early nasal CPAP-failure (32%) in our study, mostly due to RDS (89%) (4;15;16;27). This difference may be due to the lower threshold for intubation and surfactant used at our institution. Preterm infants with RDS at our neonatal intensive care unit are intubated and ventilated as soon as the FiO_2 is $>40\%$ or $PaCO_2 >8.0$ kPa and a maximum early nasal CPAP (8 cmH₂O) is reached. Treatment with surfactant is considered when mean airway pressure $\times FiO_2 >2$. In other reports, the FiO_2 threshold for intubation was higher and varied from ≥ 0.6 to 1.0 (4;15;16;27). The threshold for intubation and surfactant treatment when RDS has been established is an important issue. Prophylactic or early surfactant treatment of infants requiring mechanical ventila-

tion is more effective than late rescue treatment (17;18;28). However, with prophylactic treatment a certain amount of (surfactant sufficient) preterm infants are over-treated. Conversely, late rescue ($\text{FiO}_2 \geq 0.6$) with surfactant in an infant who is quickly deteriorating because of RDS may be disadvantageous and deleterious (29;30). Another explanation for the higher incidence of early nasal CPAP-failure in this study is the starting point of CPAP. Even if the lung is opened in the delivery room, with or without the help of mask and bag ventilation, a noncompliant surfactant-deficient lung has a tendency to collapse and lung volume is not maintained if CPAP/PEEP is not given directly to keep the lung open. In most infants in our study, early nasal CPAP was only started after transportation to the neonatal intensive care unit, which is adjacent to the delivery room, allowing the lung to collapse.

We are aware that the retrospective nature of this case-control study forms an important limitation. A selection-bias may have occurred in the delivery room when the decision was made to intubate or not and infants in the DRI-group may represent the sickest neonates with more severe RDS and higher risk for BPD. Although we found no significant differences in pulmonary clinical course parameters between the ECF and DRI groups, lack of significance can be explained by the small sample size of the study.

In summary, this study suggests that a trial of early nasal CPAP at birth is not detrimental and may be justified in case of RDS, providing early surfactant rescue is given if the infant needs to be intubated and ventilated. Based on this study, randomized controlled trials are warranted to test the hypothesis that early respiratory management of preterm neonates with RDS plays an important role in the development of BPD. Recently published or started randomized trials of early nasal CPAP are few (31;32). In 2005 we started a randomized controlled trial (EFURCI-trial, Early Functional Residual Capacity Intervention, ISRCTN 12757724) to investigate in preterm infants ≤ 32 weeks whether sustained inflation with a mechanical device (Neopuff®) and ENCPAP will be a more effective management strategy in very preterm neonates than conventional intervention and will reduce the requirement for mechanical ventilation and surfactant treatment.

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Reference List

1. The International Liaison Committee on Resuscitation (ILCOR) consensus on science with treatment recommendations for pediatric and neonatal patients: neonatal resuscitation. *Pediatrics* 2006 May;117(5):e978-e988.
2. Donn SM, Sinha SK. Can mechanical ventilation strategies reduce chronic lung disease? *Semin Neonatol* 2003 Dec;8(6):441-8.
3. Van Marter LJ, Allred EN, Pagano M, Sanocka U, Parad R, Moore M, et al. Do clinical markers of barotrauma and oxygen toxicity explain interhospital variation in rates of chronic lung disease? The Neonatology Committee for the Developmental Network. *Pediatrics* 2000 Jun;105(6):1194-201.
4. Aly H, Milner JD, Patel K, El Mohandes AA. Does the experience with the use of nasal continuous positive airway pressure improve over time in extremely low birth weight infants? *Pediatrics* 2004 Sep;114(3):697-702.
5. Avery ME, Tooley WH, Keller JB, Hurd SS, Bryan MH, Cotton RB, et al. Is chronic lung disease in low birth weight infants preventable? A survey of eight centers. *Pediatrics* 1987 Jan;79(1):26-30.
6. Lindner W, Vossbeck S, Hummler H, Pohlandt F. Delivery room management of extremely low birth weight infants: spontaneous breathing or intubation? *Pediatrics* 1999 May;103(5 Pt 1):961-7.
7. Jobe AH, Kramer BW, Moss TJ, Newnham JP, Ikegami M. Decreased indicators of lung injury with continuous positive expiratory pressure in preterm lambs. *Pediatr Res* 2002 Sep;52(3):387-92.
8. Thomson MA, Yoder BA, Winter VT, Martin H, Catland D, Siler-Khodr TM, et al. Treatment of immature baboons for 28 days with early nasal continuous positive airway pressure. *Am J Respir Crit Care Med* 2004 May 1;169(9):1054-62.
9. Bjorklund LJ, Ingimarsson J, Curstedt T, John J, Robertson B, Werner O, et al. Manual ventilation with a few large breaths at birth compromises the therapeutic effect of subsequent surfactant replacement in immature lambs. *Pediatr Res* 1997 Sep;42(3):348-55.
10. Bjorklund LJ, Ingimarsson J, Curstedt T, Larsson A, Robertson B, Werner O. Lung recruitment at birth does not improve lung function in immature lambs receiving surfactant. *Acta Anaesthesiol Scand* 2001 Sep;45(8):986-93.
11. Ingimarsson J, Bjorklund LJ, Curstedt T, Gudmundsson S, Larsson A, Robertson B, et al. Incomplete protection by prophylactic surfactant against the adverse effects of large lung inflations at birth in immature lambs. *Intensive Care Med* 2004 Jul;30(7):1446-53.
12. Ikegami M, Kallapur S, Michna J, Jobe AH. Lung injury and surfactant metabolism after hyperventilation of premature lambs. *Pediatr Res* 2000 Mar;47(3):398-404.
13. Wada K, Jobe AH, Ikegami M. Tidal volume effects on surfactant treatment responses with the initiation of ventilation in preterm lambs. *J Appl Physiol* 1997 Oct;83(4):1054-61.
14. Thomson MA. Continuous positive airway pressure and surfactant; combined data from animal experiments and clinical trials. *Biol Neonate* 2002;81 Suppl 1:16-9.
15. Ammari A, Suri M, Milisavljevic V, Sahni R, Bateman D, Sanocka U, et al. Variables Associated with the Early Failure of Nasal CPAP in Very Low Birth Weight Infants. *J Pediatr* 2005 Sep;147(3):341-7.
16. Aly H, Massaro AN, Patel K, El Mohandes AA. Is it safer to intubate premature infants in the delivery room? *Pediatrics* 2005 Jun;115(6):1660-5.
17. Soll RF, Morley CJ. Prophylactic versus selective use of surfactant in preventing morbidity and mortality in preterm infants. *Cochrane Database Syst Rev* 2001;(2):CD000510.
18. Yost CC, Soll RF. Early versus delayed selective surfactant treatment for neonatal respiratory distress syndrome. *Cochrane Database Syst Rev* 2000;(2):CD001456.
19. Jobe AH, Bancalari E. Bronchopulmonary dysplasia. *Am J Respir Crit Care Med* 2001 Jun;163(7):1723-9.
20. Jobe AH. Transition/adaptation in the delivery room and less RDS: "Don't just do something, stand there!". *J Pediatr* 2005 Sep;147(3):284-6.

21. te Pas AB, Walther FJ. Ventilation of very preterm infants in the delivery room. *Current Pediatric Reviews* 2006;2(3):187-97.
22. Subramaniam P, Henderson-Smart DJ, Davis PG. Prophylactic nasal continuous positive airways pressure for preventing morbidity and mortality in very preterm infants. *Cochrane Database Syst Rev* 2000;(2): CD001243.
23. Hussey SG, Ryan CA, Murphy BP. Comparison of three manual ventilation devices using an intubated mannequin. *Arch Dis Child Fetal Neonatal Ed* 2004 Nov;89(6):F490-F493.
24. O'donnell CP, Davis PG, Lau R, Dargaville PA, Doyle LW, Morley CJ. Neonatal resuscitation 3: Manometer use in a model of face mask ventilation. *Arch Dis Child Fetal Neonatal Ed* 2005 May;90(5):F397-400
25. Heldt GP, McIlroy MB. Dynamics of chest wall in preterm infants. *J Appl Physiol* 1987 Jan;62(1):170-4.
26. Gerhardt T, Bancalari E. Chestwall compliance in full-term and premature infants. *Acta Paediatr Scand* 1980 May;69(3):359-64.
27. Gittermann MK, Fusch C, Gittermann AR, Regazzoni BM, Moessinger AC. Early nasal continuous positive airway pressure treatment reduces the need for intubation in very low birth weight infants. *Eur J Pediatr* 1997 May;156(5):384-8.
28. Stevens TP, Blennow M, Soll RF. Early surfactant administration with brief ventilation vs selective surfactant and continued mechanical ventilation for preterm infants with or at risk for RDS. *Cochrane Database Syst Rev* 2002;(2):CD003063.
29. Bevilacqua G, Halliday H, Parmigiani S, Robertson B. Randomized multicentre trial of treatment with porcine natural surfactant for moderately severe neonatal respiratory distress syndrome. The Collaborative European Multicentre Study Group. *J Perinat Med* 1993;21(5):329-40.
30. Verder H, Albertsen P, Ebbesen F, Greisen G, Robertson B, Bertelsen A, et al. Nasal continuous positive airway pressure and early surfactant therapy for respiratory distress syndrome in newborns of less than 30 weeks' gestation. *Pediatrics* 1999 Feb;103(2):E24.
31. Finer NN, Carlo WA, Duara S, Fanaroff AA, Donovan EF, Wright LL, et al. Delivery room continuous positive airway pressure/positive end-expiratory pressure in extremely low birth weight infants: a feasibility trial. *Pediatrics* 2004 Sep;114(3):651-7.
32. Morley CJ, Davis P, Doyle L. Continuous positive airway pressure: randomized, controlled trial in Australia. *Pediatrics* 2001 Dec;108(6):1383.

