



A Review on Non-invasive Respiratory Support for Management of Respiratory Distress in Extremely Preterm Infants

Yuan Shi¹, Hemananda Muniraman², Manoj Biniwale³ and Rangasamy Ramanathan^{4*}

¹ Ministry of Education Key Laboratory of Child Development and Disorders, Key Laboratory of Pediatrics, Children's Hospital of Chongqing Medical University, Chongqing, China, ² Department of Pediatrics, Creighton School of Medicine, Omaha, NE, United States, ³ Neonatology Association Limited, Obstetrix Medical Group of Phoenix, Mednax, Arizona, AZ, United States, ⁴ Division of Neonatology, LAC+USC Medical Center, Keck School of Medicine of the University of Southern California, Los Angeles, CA, United States

OPEN ACCESS

Edited by:

Charles Christoph Roehr,
University of Oxford, United Kingdom

Reviewed by:

Aakash Pandita,
Sanjay Gandhi Post Graduate Institute
of Medical Sciences, India

MaryAnn Volpe,
Tufts University School of Medicine,
United States

*Correspondence:

Rangasamy Ramanathan
ramanath@usc.edu

Specialty section:

This article was submitted to
Neonatology,
a section of the journal
Frontiers in Pediatrics

Received: 07 December 2019

Accepted: 29 April 2020

Published: 28 May 2020

Citation:

Shi Y, Muniraman H, Biniwale M and
Ramanathan R (2020) A Review on
Non-invasive Respiratory Support for
Management of Respiratory Distress
in Extremely Preterm Infants.
Front. Pediatr. 8:270.
doi: 10.3389/fped.2020.00270

Majority of extremely preterm infants require positive pressure ventilatory support at the time of delivery or during the transitional period. Most of these infants present with respiratory distress (RD) and continue to require significant respiratory support in the neonatal intensive care unit (NICU). Bronchopulmonary dysplasia (BPD) remains as one of the major morbidities among survivors of the extremely preterm infants. BPD is associated with long-term adverse pulmonary and neurological outcomes. Invasive mechanical ventilation (IMV) and supplemental oxygen are two major risk factors for the development of BPD. Non-invasive ventilation (NIV) has been shown to decrease the need for IMV and reduce the risk of BPD when compared to IMV. This article reviews respiratory management with current NIV support strategies in extremely preterm infants both in delivery room as well as in the NICU and discusses the evidence to support commonly used NIV modes including nasal continuous positive airway pressure (NCPAP), nasal intermittent positive pressure ventilation (NIPPV), bi-level positive pressure (Bi-PAP), high flow nasal cannula (HFNC), and newer NIV strategies currently being studied including, nasal high frequency ventilation (NHFV) and non-invasive neutrally adjusted ventilatory assist (NIV-NAVA). Randomized, clinical trials have shown that early NIPPV is superior to NCPAP to decrease the need for intubation and IMV in preterm infants with RD. It is also important to understand that selection of the device used to deliver NIPPV has a significant impact on its success. Ventilator generated NIPPV results in significantly lower rates of extubation failures when compared to Bi-PAP. Future studies should address synchronized NIPPV including NIV-NAVA and early rescue use of NHFV in the respiratory management of extremely preterm infants.

Keywords: bronchopulmonary dysplasia (BPD), nasal continuous positive airway pressure (NCPAP), nasal intermittent positive pressure ventilation (NIPPV), high flow nasal cannula (HFNC), nasal high frequency ventilation (NHFV), noninvasive ventilation (NIV), noninvasive ventilation-neurally adjusted ventilatory assist (NIVNAVA)

INTRODUCTION

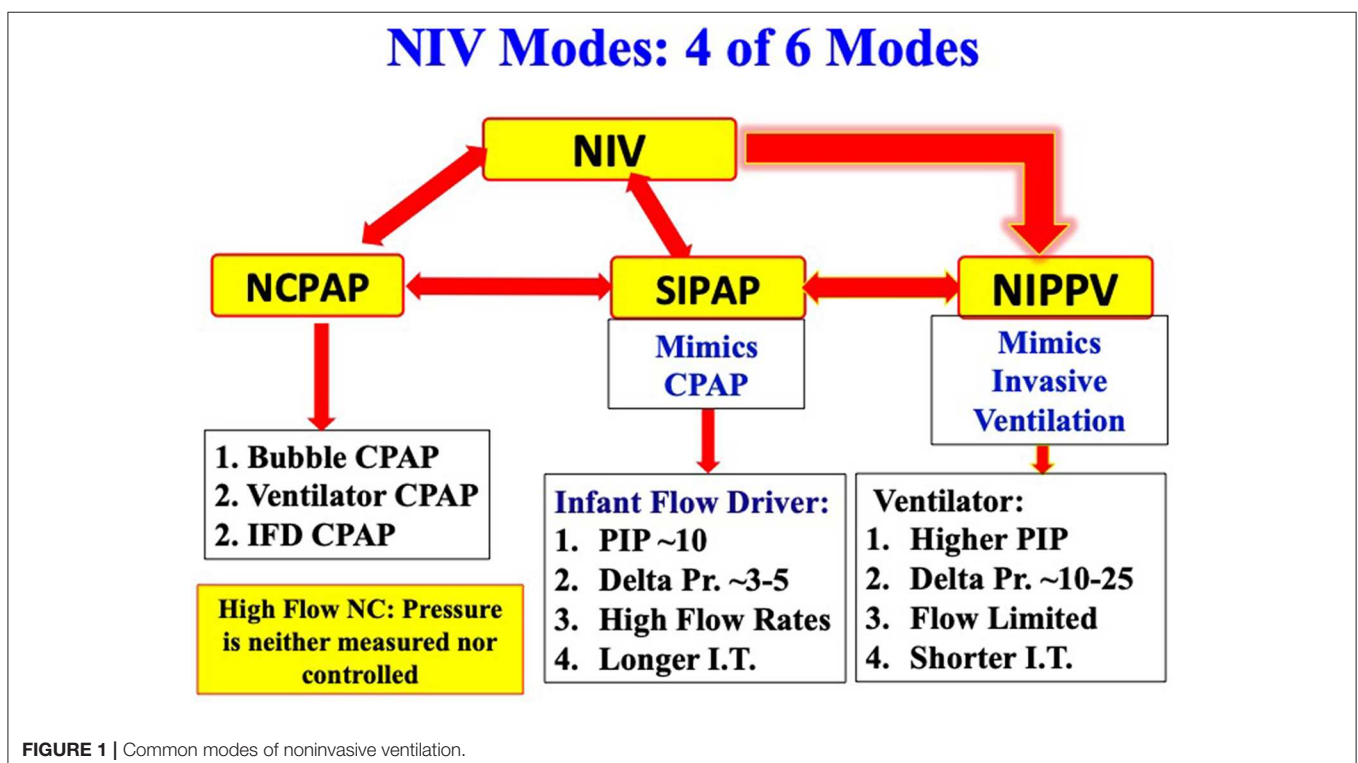
Providing optimal ventilation strategies remains the key to success of managing extremely preterm infants. Majority of the extremely preterm infants have respiratory distress (RD) needing significant respiratory support immediately after birth or after admission to the neonatal intensive care unit (NICU) due to poor inspiratory effort, weak intercostal muscles, and poor diaphragmatic function. These infants are at very high risk of developing bronchopulmonary dysplasia (BPD) and adverse neurodevelopmental outcomes, which are directly related to the duration of invasive mechanical ventilation (IMV) and supplemental oxygen. The strong association between ventilator dependency and neurologic injury, such as severe intraventricular hemorrhage (IVH) and periventricular leucomalacia, emphasizes the severity of their illness (1). A recent study showed more than 60 days of positive pressure support regardless of invasive or non-invasive ventilation (NIV) mode was associated with a higher risk for neurodevelopmental problems (2). Avoiding intubation and using NIV modes in preterm infants minimizes the risk for lung injury and optimizes neonatal outcomes. Use of nasal continuous positive airway pressure (NCPAP) in the delivery room and nasal intermittent positive pressure ventilation (NIPPV) in the NICU has been shown to decrease the need for IMV in extremely preterm infants without increasing major morbidities (3).

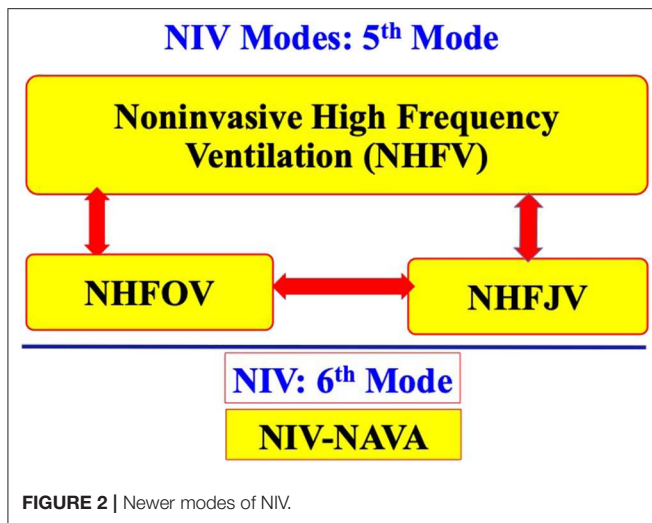
In extremely preterm infants, BPD is associated with long term impaired pulmonary function and adverse neurological outcomes (4–6). While the etiology of BPD is multifactorial, lung injury particularly with IMV and resulting inflammation play

a major role in the pathogenesis (7). NIV has been shown to reduce the risk of BPD when compared to IMV (8). Ventilatory practices have evolved over the last few decades with preference for NIV in the management of respiratory distress syndrome (RDS) in extremely preterm infants (4). Also, the definition of BPD is constantly evolving. Existing definitions mostly relied on level and duration of supplemental oxygen and did not take into account the major changes in NIV modes that are currently used in preterm infants. Recently, in a study using 18 pre-specified definitions of BPD that used disease severity based on level of respiratory support and supplemental oxygen at 36 weeks' postmenstrual age (PMA), only mode of respiratory support best predicted early childhood morbidity, regardless of supplemental oxygen use. Our focus should be to assess and follow the extremely preterm infants based on level of non-invasive as well as invasive positive pressure support at the time of discharge and after discharge (9).

MODES OF NON-INVASIVE VENTILATION

Six modes of NIV are currently used in extremely preterm infants. Four of the six modes commonly used in most of the NICUs include NCPAP, bilevel positive airway pressure (Bi-PAP) or sigh breaths above a baseline CPAP pressure (Si-PAP), NIPPV and high flow nasal cannula (HFNC) (**Figure 1**). Remaining 2 modes, namely, nasal high frequency ventilation (NHFV) using nasal high frequency oscillatory ventilation (NHFOV) or nasal high frequency jet ventilation (NHFJV) and non-invasive neurally adjusted ventilatory assist (NIV-NAVA) are





not well-studied in extremely preterm infants and need further evaluations before routine use of these modes (Figure 2). Positive pressure delivery during NIV involves pressure generators, nasal interfaces (Figure 3) and ability to provide one or two levels of pressures at different ventilator rates. NIV in the NICU can be provided by these modalities either as a primary mode of respiratory support or following extubation after a period of IMV. NHFV modes are also being used increasingly as a rescue mode to treat hypercarbia and to decrease the need for intubation in extremely preterm infants failing other modes of NIV support. NIV can also be used in combination with early, rescue surfactant treatment. Surfactant can be delivered using invasive techniques such as INSURE (INTubation, SURfactant, and Extubation), minimally invasive techniques including SurE (surfactant without endotracheal intubation) using a feeding tube or a specially designed catheter, laryngeal mask airway, or non-invasive technique, like, nebulization (10–13).

MECHANISMS OF NIV

Physiological mechanisms leading to improvement in the lung mechanics are somewhat similar with all these modes of NIV. NCPAP reduces upper airway resistance, helps to establish functional residual capacity (FRC), decreases chest wall distortion, augments spontaneous breathing efforts, preserves endogenous surfactant, decreases the need for surfactant administration, and decreases the need for, and duration of IMV (14). However, in patients with hypopnea or apnea, NCPAP often fails, needing intubation and IMV. Providing a backup rate using NIPPV with adequate peak inspiratory pressure (PIP) decreases apneic spells, improves ventilation and decreases the need for intubation. NIPPV is a time cycled, pressure limited mode of ventilation. Conventional ventilator is used to generate two levels of pressures, namely, PIP and positive end expiratory pressure (PEEP). Additionally, a backup rate is provided typically using longer inspiratory time. Benefits of NIPPV mode include all of the benefits of NCPAP listed above,

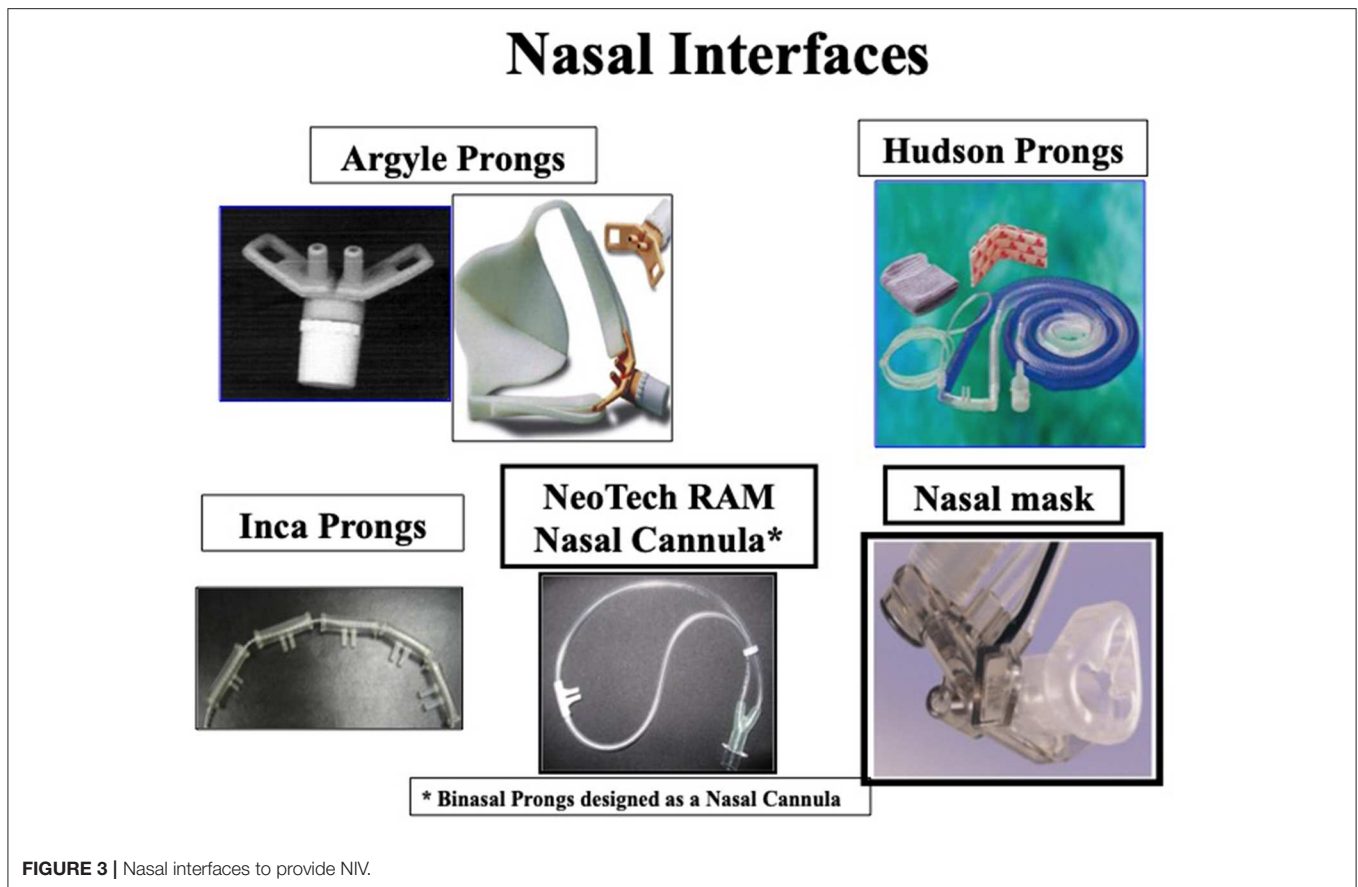
and pharyngeal dilation with further decrease in upper airway resistance, augmentation of spontaneous inspiratory effort via Head's paradoxical reflex, improving compliance and reopening of partially collapsed airways, increase in FRC, increase in tidal volume (V_t) and minute volume, better alveolar recruitment due to higher mean airway pressure (MAP), reduction in chest wall distortion, and improved respiratory unloading with decrease in work of breathing (15). Head's paradoxical reflex is seen typically during a rapid inflation of the lungs causing a deep inspiration or gasp. It is mediated by the irritant receptors of the major airways receptive to lung inflation. The reflex is seen most commonly on the first day and may help to establish and maintain FRC. Head's paradoxical reflex has also shown to possibly increase neural inspiratory time in patients receiving NIV-NAVA (16). NIPPV also stimulates the Hering-Breuer inflation reflex with inflation of lungs resulting in cessation of respiratory activity preventing hyperinflation. This reflex is mediated through the stretch receptors in the smooth muscles of the major airways and is time-dependent with a longer inspiratory time resulting in a longer period of respiratory inhibition before the next breath. Both NCPAP and NIPPV modes may trigger this reflex, causing slower spontaneous respiratory rate. In the preterm infants, this reflex produces rapid, shallow tidal breathing. In older infants this reflex prevents excessive tidal volumes and can only be stimulated if the inflating volume is increased beyond a critical threshold.

Proposed mechanisms for the use of HFNC include washout of nasopharyngeal dead space, decrease in inspiratory resistance, and provision of positive pressure. However, pressure generated during HFNC is neither measured nor controlled by the clinician and is very unpredictable (17). NHFV modes promote better lung recruitment and removal of carbon dioxide.

DEVICES FOR DELIVERING NIV

NCPAP may be provided using a water column as a resistor to generate CPAP (Bubble CPAP), or using a flow generator as in infant flow driver (IFD) device, or using a conventional ventilator with continuous or variable flow rates. Studies comparing different modes of providing NCPAP have shown no significant difference in extubation failure rates (18, 19). Most commonly used bi-level modes include Bi-PAP, Si-PAP, and Duo-PAP. IFDs are variable flow devices, and generate two levels of pressures, a high pressure and a low pressure or CPAP by varying the flow rates. Bi-PAP/Si-PAP mimics NCPAP due to low delta pressure in these modes. The delta pressure during bi-level mode is between 5 and 10 cmH_2O , which often is not enough to treat hypercapnia or support poor spontaneous respiratory efforts.

Most of the conventional ventilators can provide NIPPV with PIP, PEEP, and rate to provide adequate support. Furthermore, flow can be adjusted to provide adequate pressure or compensate for leaks. Newer ventilators have NIV modes with excellent leak compensation. Only one type of ventilator provides NIV-NAVA mode. Both high frequency oscillatory ventilator as well as high frequency jet ventilators can be used to provide NHFV. HFNC



is provided by dedicated devices where only flow is adjusted to optimize gas exchange.

NASAL INTERFACES IN DELIVERY ROOM AND NICU

Common interfaces used in the delivery room (DR) to provide NIV support are round and anatomical mask, single or bi-nasal nasopharyngeal prongs, bi-nasal prongs, nasal mask, or RAM nasal cannula (NC) (Neotech RAM Nasal Cannula[®], Neotech Products, Valencia, California, USA) with pressure generating devices including self-inflating or flow-inflating bag, and T piece resuscitator (20). Bag and mask resuscitation is often not effective in the DR even when performed by the experienced personnel in extremely preterm infants. Three major issues with bag and mask ventilation are: mask leak, upper airway obstruction from the tongue falling backwards toward the oropharynx and increase in dead space with the gas in the oropharynx not contributing to gas exchange (21–24). Corrective ventilation steps during resuscitation are taught using MRSOPA mnemonic: Mask adjustment, Repositioning airway, Suctioning, Opening the mouth, Increasing inspiratory pressure, and Alternative airway. These steps are not always successful, especially among very preterm infants. The greater the number of MRSOPA steps used in the DR, the more likely intubation occurred (25). In another

study, MRSOPA maneuvers improved tidal volume delivery in some cases, but, worsened exhaled tidal volumes in others. In fact, these authors found MRSOPA steps actually induced mask leak and airway obstruction in some cases (26). In a randomized, controlled trial, use of NC vs. face mask for primary neonatal resuscitation in the DR in more mature neonates (mean GA 36 weeks), NC use resulted in significantly less need for intubation (0.6 vs. 6.3%; $p < 0.001$) and chest compressions (1.65 vs. 8.28%; $p = 0.001$) in the NC group (27).

Successful use of RAM NC for the resuscitation of very low birth weight infants and decreased the need for intubation even among the lower gestational age infants (mean GA 27 weeks) has been reported (28). A recent study in <29 weeks' gestation infants used sustained lung inflation (SLI) followed by NCPAP ranging from 6 to 8 cmH₂O, using RAM NC resulted in a significant reduction in intubation rates in the DR (29). Success with the use of RAM NC as an interface is likely due to delivery of tidal volume through the nasopharynx, eliminating dead space by avoiding oropharyngeal space, ease of application with T-Piece resuscitator, and minimizing upper airway obstruction (Figure 4) (30). In addition, sustained inflation or NIPPV may be applied in preterm infants needing additional support without manipulating the tiny infants during resuscitation. Among the pressure delivery devices, T piece resuscitator delivers targeted inflation pressure more consistently compared to self-inflating or flow inflating

Advantages of RAM NC over Bag & Mask in the Delivery Room

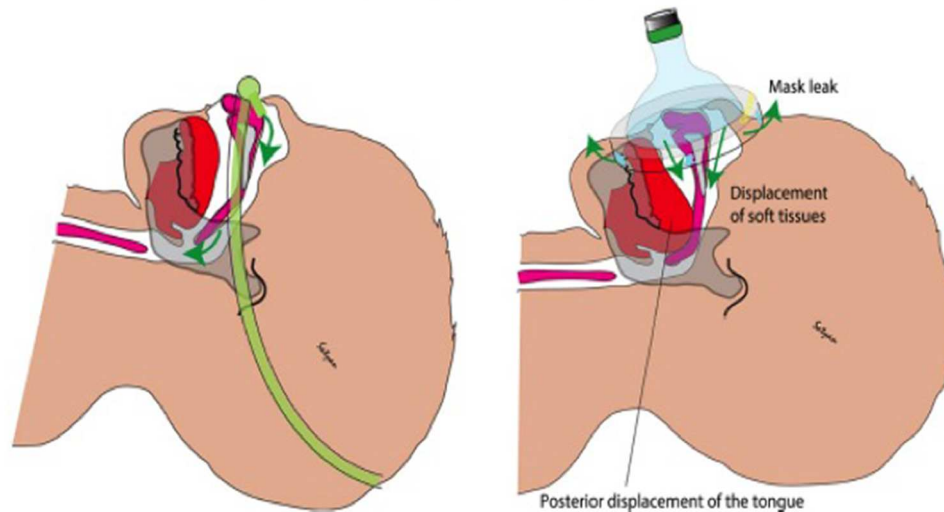


Fig. 17.10 Advantages of RAM Cannula Over Mask Ventilation In the DR. Copyright: Satyan Lakshminrusimha.

No leaks; By-passes oro-pharynx and decreases dead space; easy to use

Modified from Essentials of Neonatal Ventilation, 1st edition, Elsevier; copyright Satyan Lakshminrusimha (used with permission)

FIGURE 4 | Application of RAM NC in the NICU.

bag although this has not been shown to improve clinical outcomes (31).

Various interfaces used in the NICU include varieties of short nasal prongs, RAM NC and nasal masks. There is considerable variation in measured resistance between these interfaces. When applying smallest size interfaces for extremely preterm infants all nasal interfaces result in decrease in pressures due to high resistance. Pressure drop may vary based on the set flow, internal diameter and length of the prongs. Use of interfaces with high resistance may result in a greater drop in delivered airway pressure in comparison to set pressure (32). Application and advantages of RAM NC to provide NIV in the NICU are shown in **Figure 5**. Clinicians need to be aware of adjusting pressure and flow settings while using the ventilator to provide NIV. A meta-analysis of studies with nasal mask comparing to binasal prongs showed significantly decreased the risk of CPAP failure (4 RCTs [$N = 459$]; relative risk [RR]: 0.63; 95% confidence interval [CI]: 0.45–0.88; $P = 0.007$; $I^2 = 0\%$, NNT: 9), and the incidence of moderate to severe nasal trauma (3 RCTs [$N = 275$], RR: 0.41; 95%CI, 0.24–0.72; $P = 0.002$; $I^2 = 74\%$, NNT: 6) (33). Larger studies are needed to validate safety and efficacy of using nasal masks in extremely preterm infants.

Pressure Transmission During NIPPV

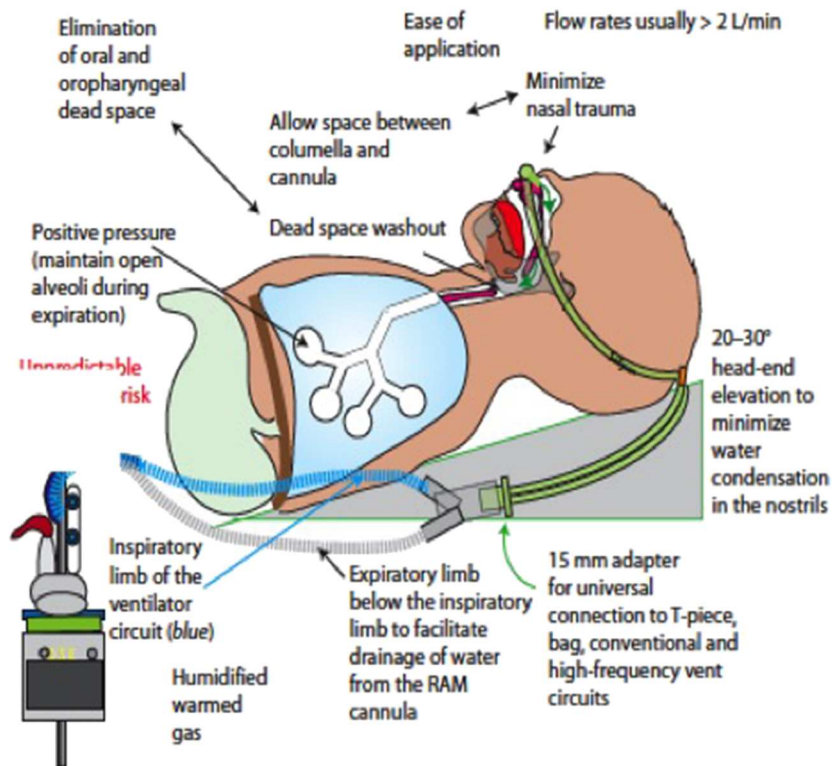
Pressure transmission to the hypopharynx or to the lung during NIPPV is difficult to measure. It depends on the size of the

prongs, length of tubing, leaks around the nostrils, and set inspiratory time and whether the infant's mouth is open or closed. Using a computerized test lung simulator, pressure transmission using different size RAM NCs has been reported. In this well-designed study with a 30% leak, authors demonstrated around 70, 80, and 90% of set PIP delivered to the hypopharynx with preemie, newborn, and infant size RAM NCs respectively (34). More studies are needed comparing similar diameter prongs and similar leak settings. If a higher CPAP or PEEP is needed, then, the leak at the nasal interface may be decreased by using cannulaide[®] (Beever Medical Solutions, OR, USA).

Temperature and Humidity During NIV

Heating and humidification of inspired gas is a routine practice when providing respiratory support in infants. Unconditioned dry and cold gas can result in impaired ciliary function, reduced clearance of secretions, damage to the airway mucosa which may impair lung function by reducing compliance and FRC. Different NIV interfaces deliver inspiratory gases of variable temperature and humidity. Some HFNC and variable flow CPAP devices at higher gas flow may not achieve the recommended temperature and humidity (35). Higher NHFOV settings with low frequencies, high amplitudes, and high inspiratory to expiratory ratios may also place infants at an increased risk of upper airway injury due to decreased humidification (36). There are no studies on humidity or temperature of gas

Use of RAM Nasal Cannula in the NICU



Modified from *Essentials Neonatal Ventilation*, 1st edition, Elsevier; copyright Satyan Lakshminrusimha (used with permission)

FIGURE 5 | NCPAP in delivery room and early failures.

delivered in the pharynx with RAM NC or other binasal prongs. Humidification is evidenced by condensation in the tubing of RAM NC. Temperature measurement is done at the wye and beyond that, the tubing length is 11 cm to the prongs. With continuous flow of heated and humidified gas, a significant drop in temperature between the wye and the patient's nasal interface is not expected. Inspiratory gas also gets heated and humidified by the patient's nasopharynx.

NIV FOR STABILIZATION IN THE DELIVERY ROOM

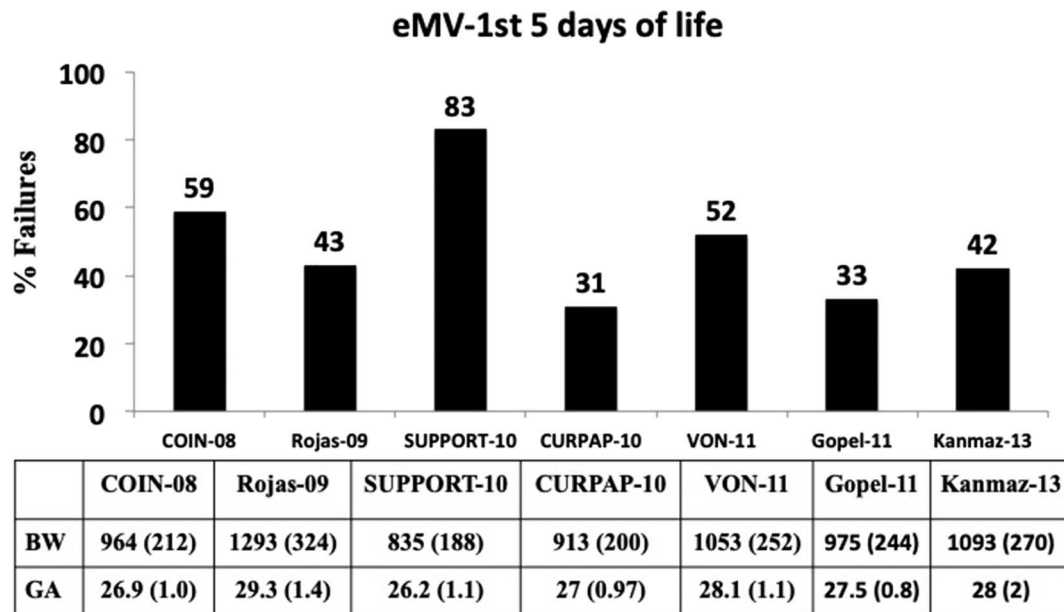
Establishment of FRC during the isovolumic transformation of a fluid filled lung to an air breathing lung is critical for successful adaptation and post-natal transition (20). Extremely preterm infants are at high risk of respiratory distress and maladaptation owing to immature lungs, insufficient production of surfactant, highly compliant chest wall and immature respiratory center control (37). As a result, more than 70% of extremely preterm infants require positive pressure support (38). Strategies to optimize lung recruitment and establishing FRC at this crucial

period can play a major role in decreasing respiratory morbidities and mortality in this vulnerable population. Until early 2000s, elective intubation with prophylactic administration of surfactant was the standard of care in the initial management of extremely preterm infants. After large clinical trials showing benefits of NCPAP use in the DR to decrease need for intubation and IMV (39, 40) use of NIV during stabilization and initial treatment of respiratory distress has significantly increased (4).

NCPAP IN DELIVERY ROOM

CPAP has been shown to be effective in establishing FRC (37). Two large randomized controlled trials (RCT) compared NCPAP with routine intubation in the delivery room (39, 40). The CPAP or intubation at birth (COIN) trial randomized 610 spontaneously breathing infants born at 25 to 28 weeks' gestation with signs of respiratory distress at 5 min of life to receive either CPAP or endotracheal intubation. Infants intubated due to respiratory distress before 5 min of age were excluded. NCPAP of 8 cmH₂O was used in this study. There was no difference in the primary outcome, namely, death or BPD

NCPAP in the Delivery Room & Early Failures (eMV in the 1st <5 days) 2008-2013



Modified from Fischer HS, Buhner C. *Pediatrics* 132:e1351-e60;2013

FIGURE 6 | Advantages of RAM NC over Bag & Mask in the delivery room.

between the two groups; however, there was a higher incidence of pneumothorax in the CPAP group (39). The surfactant positive airway pressure and pulse oximetry (SUPPORT) trial from USA randomized 1,316 infants between 24 and 28 weeks' gestational age to receive NCPAP or endotracheal intubation along with administration of surfactant. Overall mortality (47.8 and 51%, respectively) and BPD rates were similar between the NCPAP and the intubation with surfactant group (40). Evidence from these studies showed that NCPAP was as effective as routine intubation in the extremely preterm infants. To date, seven RCTs using NCPAP in the delivery room have been published. Failure rates needing intubation and IMV ranged from 31 to 83% (**Figure 6**; **Table 1**) (39–45, 47). None of individual clinical trials showed benefit in the primary outcome, namely, death or BPD. However, systematic review and meta-analysis of these studies showed a small but significant benefit in decreasing death or BPD, with a number needed to treat (NNT) of 25 (48). In a long term follow up study by Doyle et al. despite substantial increase in the use of NCPAP, there was no decrease in BPD and more importantly, no improvement in lung function was seen at 8 years of age (49). Exact reasons for the lack of benefit with NCPAP are not clear. It may be due to inability to recruit lungs with inadequate CPAP pressures due to leaks with nasal interfaces or lack of augmentation of breaths as provided during NIPPV.

NIPPV IN THE DELIVERY ROOM

There are no randomized clinical trials of using NIPPV compared to CPAP in the DR. Infants are typically placed on either CPAP or NIPPV after initial resuscitation. In a retrospective study in very low birth weight infants, comparing positive pressure ventilation (PPV) using a face mask to directly placing on NIPPV with RAM nasal cannula at birth, NIPPV use was associated with a significantly decreased need for intubation in the DR (31 vs. 85%) including among the extremely preterm infants born at 24–27 weeks of gestation, decreased need for chest compressions (11 vs. 31%), and decreased the need for IMV at 24 h of age (38 vs. 66%) (28).

USE OF SUSTAINED LUNG INFLATION IN DELIVERY ROOM

Sustained lung inflation (SLI) strategy may result in better lung recruitment immediately after birth through delivery of a PIP of 15–30 cmH₂O for a sustained period of time, typically, 10–15 s to the infant airways via a nasopharyngeal tube or mask or NC, followed by CPAP. SLI procedure creates a transepithelial pressure gradient across the alveolar-capillary membrane and helps to move fluid from the alveoli into the interstitial space and subsequent removal of this fluid via lung lymphatics

TABLE 1 | Studies comparing NCPAP and intubation with IMV.

References	Intervention (n)	GA, weeks	BPD (%)	Death (%)	Combined BPD and death (%)	Intubation rates in NCPAP group (%)
Morley et al. (39)	NCPAP (307) vs. IMV (303)	25–28	29 vs. 35	6.5 vs. 5.9	34 vs. 39	59
SUPPORT et al. (40)	NCPAP (663) vs. IMV (653)	24–28	40 vs. 44	14 vs. 17	49 vs. 54	83
Dunn et al. (41)	NCPAP (223) vs. INSURE (216) vs. IMV (209)	26–29	n/a	4 vs. 7 vs 7	30 vs. 28 vs. 36	52
Fojas et al. (42)	NCPAP (137) vs. INSURE (141)	27–30	59 vs. 49	9 vs. 9	62 vs. 54	53
Sandri et al. (43)	NCPAP (105) vs. INSURE (103)	25–29	n/a	n/a	21 vs. 22	31
Göpel et al. (44)	NCPAP ± LISA (108) vs. nCPAP ± INSURE (112)	26–28	8 vs. 13	n/a	14 vs. 15	46
Kanmaz et al. (45)	NCPAP + LISA (100) vs. nCPAP INSURE (100)	<30	10 vs. 20*	16 vs. 13	34 vs. 45	40
Tapia et al. (46)	NCPAP + INSURE (131) vs. MV (125)	800–1,500g	7 vs. 10	8 vs. 9	14 vs. 19	30

NCPAP, Nasal continuous positive pressure ventilation; IMV, Invasive mechanical ventilation; INSURE, Intubation, SURfactant and Extubation; LISA, Less invasive surfactant administration; GA, Gestational age (weeks) *P < 0.05.

and pulmonary microcirculation. SLI superimposed on PEEP may have beneficial effects, like, maintaining adequate FRC, promoting optimal gas exchange, improving lung mechanics, and reducing the need for intubation in the DR (50). A recent large multicenter study in extremely preterm infants requiring resuscitation at birth, a ventilation strategy involving 2 SLIs at maximal PIP of 25 cmH₂O for 15 s, compared with standard intermittent positive pressure ventilation, did not reduce the risk of BPD or death at 36 weeks postmenstrual age. The study was stopped early due to safety concerns with increased rates of death in the infants receiving SLI (51). Reasons for increased mortality with SLI in this study are not clear. Updated systematic review and meta-analysis of SLI vs. intermittent positive pressure ventilation and continuous positive airway pressure for the prevention of hospital mortality and morbidity in preterm infants showed no difference in the risk of the primary outcome of death in the delivery room or before hospital discharge, although SLI was associated with increased risk of death in the first 2 days after birth, with no evidence of efficacy for SLI prevent other neonatal morbidities. Duration of mechanical ventilation was shorter in the SLI group but did not translate into better long term pulmonary outcomes. These findings do not support the routine use of SLI in preterm infants at birth in the delivery room (52, 53).

HIGH FLOW NASAL CANNULA FOR STABILIZATION IN THE DELIVERY ROOM

Only one study had evaluated HFNC during stabilization at birth. Reynolds et al. performed a pilot study evaluating use of HFNC (6–7 LPM) in stabilizing infants <30 weeks' gestation; 25 of 28 infants were successfully stabilized with HFNC, 48% of the infants received surfactant and 60% of the infants remained on

HFNC at 72 h of age (54). They concluded that it is feasible to use HFNC in preterm infants. Additional studies in extremely preterm infants are needed.

EARLY USE OF NIV IN NICU

The major risk factors for BPD in extremely infants are treatment with oxygen and IMV. A meta-analysis of 7 randomized controlled trials including 3,289 patients showed that avoiding IMV reduced the combined outcome of death or BPD in preterm infants <30 weeks' gestational age (47). At present, NCPAP and NIPPV are the two most common modes used either as a primary mode or rescue mode of NIV support in the NICU.

EARLY NCPAP

Previous systematic reviews of randomized clinical trials of preterm infants found that the early use of NCPAP to avoid IMV decreased BPD, death, or both compared with the respiratory management using routine intubation (48, 55). However, one major disadvantage of using only NCPAP without intubation is a delay in the administering surfactant that is generally given via an endotracheal tube after intubation. A large multicenter study comparing early NCPAP to intubation and surfactant within 1 h of age in infants less than 28 weeks of gestation did not show any significant differences in long term morbidities (40). Several factors in extremely preterm infants including gestational age <26 weeks, birth weight < 750 g, need PPV in the DR, FiO₂ >0.30 and severe RDS on chest x-ray contribute to NCPAP failures (56). CPAP failure is associated with increased risk of mortality and major morbidities, including BPD, both in infants <29 weeks' and in infants between 29 and 32 weeks' GA (57).

USE OF NCPAP WITH SURFACTANT THERAPY

Few RCTs have evaluated routine NCPAP with NCPAP after surfactant via INSURE or LISA technique (41–46). In the DR management trial, infants 26 to 29 weeks' gestation were randomized to 3 groups: prophylactic surfactant followed by a period of mechanical ventilation for at least 6h, prophylactic INSURE within 30 min followed by bubble NCPAP or initial management with bubble NCPAP and selective surfactant treatment. There were no differences in death or moderate to severe BPD (NCPAP 4.1% vs. INSURE 7% vs. prophylactic surfactant 7.2%), and in pneumothorax (5.4% vs. 3.2% vs. 4.8%) in these 3 groups (41). In another multicenter, RCT from the South American Neocosur Network, early bubble CPAP and selective surfactant by INSURE technique reduced the need for mechanical ventilation and surfactant; however, there were no differences in the rates of death or BPD (46). A multicenter RCT from Germany included 220 infants between 26 and 29 weeks' gestation and reported decreased need for mechanical ventilation in infants treated with NCPAP and surfactant administration via LISA technique, but no decrease in BPD (44). However, a similar study from Turkey reported decrease in both the need for mechanical ventilation and BPD (10 vs. 20%) when treated with NCPAP and LISA (45). A recent study showed SurE technique using a thin catheter for surfactant delivery resulted in decreased need for MV and less BPD (11). In a meta-analysis including majority of the above mentioned studies, Fischer et al. concluded that avoiding early IMV by using NCPAP with or without surfactant resulted in a small but significant beneficial effect on preventing BPD, with a number needed to treat (NNT) of 35 (47) (Table 1). Furthermore, NCPAP failures in preterm infants <29 weeks GA is associated with increase in mortality, BPD, death or BPD, and necrotizing enterocolitis (NEC) (57).

EARLY NCPAP VS. NIPPV USE IN NICU

A major reason for lack of benefit in the NCPAP trials is due to high rates of NCPAP failures, requiring intubation

within 3–7 days after randomization. Most common reasons for NCPAP failures are recurrent apnea, bradycardia or desaturation episodes, hypopnea, need for higher pressures (NCPAP > 8 cmH₂O), and/or severe respiratory acidosis. NCPAP when used as a primary mode or following a period of IMV has been shown to result in failure rates of 31% to 83% (Table 1), requiring intubation or re-intubation. NIPPV augments NCPAP and has been shown to be more effective than NCPAP after extubation and in the treatment of apnea of prematurity (58, 59).

Several large RCTs comparing early NCPAP with early NIPPV have been published (60–69) (Table 2). Of the 10 studies reviewed, 6 studies enrolled patients prior to surfactant administration (60–64, 69), 2 studies had mixed enrollment (65, 66), and 2 studies enrolled after INSURE technique (67, 68). Four of these studies reported decreased rates of IMV (64, 65, 68, 69). Three studies reported decreased respiratory failure and duration of oxygen requirement (60, 61, 67). Three studies, where rescue surfactant via INSURE and LISA was provided, showed decreased rates of BPD (64, 68, 69). A recent Cochrane meta-analysis involving 10 trials enrolling 1,061 infants showed significantly decreased rates of respiratory failure [relative risk] RR 0.61 (95% CI 0.51, 0.82)], decreased need for intubation [RR: 0.78 (95% CI 0.64, 0.94)], and NNT to prevent one extubation failure was 17 with NIPPV. There were no differences in the rates of BPD (RR: 0.78; 95% CI 0.58, 1.06) and mortality (RR: 0.77 (95% CI 0.51, 1.15); however, in one study, combining surfactant with NIPPV led to a reduction in BPD. There were no differences in pneumothorax, NEC, IVH, and retinopathy of prematurity (70).

One of the largest RCT involving 1,009 patients <1 kg at birth, comparing NCPAP with Si-PAP or NIPPV, reported no difference in extubation failures (61.8 vs. 59.5%), survival with BPD (31 vs. 33.9%), and death or BPD (36.7 vs. 38.4%) (62). In this pragmatic study, more than half of the centers used IFD device to deliver “NIPPV,” and in centers using a ventilator to deliver NIPPV, maximum PIP that could be used was limited to 18 cmH₂O. Even though, the authors described this study as NIPPV vs. NCPAP, this was truly a study comparing Si-PAP with NCPAP. Post-randomization failures needing intubation were very high, most likely, secondary to lower delta pressures used in the “NIPPV”

TABLE 2 | Studies comparing NIPPV and NCPAP.

References	n	Synchroniz-ation	Surfactant prior	GA, weeks	Intubation/respiratory failure	Death	BPD
Armanian et al. (60)	98	No	No	<35	4 vs. 2	4 vs. 2	n/a
Bisceglia et al. (61)	88	No	No	28–34	2 vs. 2	0 vs. 0	4 vs. 8
Kirplani et al. (62)	185	Some	No	<30	21 vs. 29	3 vs. 4	19 vs. 14
Meneses et al. (63)	200	No	No	26–33	58 vs. 64	22 vs. 26	26 vs. 25
Kugelman et al. (64)	84	Yes	No	24–34	25 vs. 46*	0 vs. 0	2 vs. 17*
Sai Sunil Kishore et al. (65)	76	No	Some	28–34	19 vs. 41*	13 vs. 23	3 vs. 10
Salama et al. (66)	60	Yes	Some	28–34	10 vs. 20	0 vs. 3	3 vs. 6
Lista et al. (67)	40	Yes	Yes	28–34	10 vs. 15	0 vs. 0	0 vs. 0
Ramanathan et al. (68)	110	No	Yes	26–29	17 vs. 42*	2 vs. 2	22 vs. 39
Oncel et al. (69)	200	No	No	26–32	13 vs. 29*	4 vs. 6	7 vs. 16*

NIPPV, Nasal intermittent positive pressure ventilation; NCPAP, Nasal continuous positive pressure ventilation. **p* < 0.05.

TABLE 3 | Suggested settings for CPAP, Bi-PAP, NIPPV, and HFNC.

Mode of NIV	Initial settings	Max settings	Weaning parameters	Lowest settings
NCPAP	5–6 cmH ₂ O	8–10 cmH ₂ O	1 cmH ₂ O	4 cmH ₂ O
Bi-PAP	High Pressure 10 cmH ₂ O Low Pressure 5 cmH ₂ O Rate 20/min	High Pressure 15 cmH ₂ O Low Pressure 8 cmH ₂ O Rate 30/min	1 cmH ₂ O wean the rate by 2–4 /min every 6 h	High/Low Pressure 8/5 cm H ₂ O Rate 0
NIPPV	PIP 20 cmH ₂ O PEEP 6 cmH ₂ O Inspiratory time 0.5 s Rate 40/min	PIP 35–38 cmH ₂ O PEEP 8–10 cmH ₂ O Rate 50/min	wean PIP first by 1–2 until lowest possible PIP wean the rate by 2–4 /min every 6 h	PIP 12 or 15 cmH ₂ O PEEP 4–5 cmH ₂ O Rate 20/min
HFNC gas flow	4–6 L/min	8 L/min.	0.5–1.0 L/min	1–4 L/min

NIV, non-invasive ventilation; NCPAP, nasal continuous positive airway pressure; NIPPV, nasal intermittent positive pressure ventilation; HFNC, high flow nasal cannula; PIP, peak inspiratory pressure; PEEP, positive end expiratory pressure.

group. However, Lemyre et al. included 185 infants from this study who were randomized prior to intubation and surfactant administration in their Cochrane review and found decreased need for intubation and respiratory failure (70).

Early NIPPV appears to be superior to NCPAP alone for decreasing respiratory failure and the need for intubation and IMV among preterm infants with RDS (70). Another important factor is related to the devices used to deliver NIPPV. NNT to decrease respiratory failure and intubation with a ventilator delivered NIPPV was 13 (70). Current evidence suggests early NIPPV delivered with a ventilator and minimally invasive technique for early, rescue surfactant therapy, like, LISA may be the most effective strategy to minimize IMV and improve outcomes in extremely preterm infants. Recommended settings for NIPPV, NCPAP, and HFNC are shown in the **Table 3**.

There are no studies comparing weaning strategies from NIPPV to CPAP. Several factors are to be considered while weaning including underlying pulmonary disease, intermittent hypoxic episodes, post-natal age, growth, oxygen requirement, and gas exchange. Individual patient specific weaning strategy is encouraged. For extremely preterm infants we (HM, MB, RR) typically wean PIP first before weaning the rate. When the PIP is around 12 or 15 cmH₂O, PEEP at 5–6 cmH₂O, and FiO₂ < 0.30, we wean the rate by 2–4 bpm every 6 h and transition to NCPAP. When patient is stable on NCPAP for 12–24 h, we wean to low flow NC (<2 lpm).

NIPPV VS. NCPAP POST-EXTUBATION

Infant receiving invasive ventilation are at high risk for developing complications such as increased hemodynamic instability, increased airway resistance, acute and chronic airway trauma, increased ventilation associated infections and reduced clearance of secretions. Minimizing IMV and extubating to NIV may aid in avoiding these undesirable side effects. Choice of post-extubation respiratory support is based on several factors including level of respiratory support at the time of extubation, duration of respiratory support, underlying lung pathology, and associated clinical problems as well as infant's hemodynamic status.

TABLE 4 | Interventions to improve rates of successful extubation in preterm infants.

Preventing extubation failures	Risk ratio [95% CI]	NNT [95% CI]
NCPAP vs. Head-Box	0.59 [0.48–0.72]	6 [3–9]
NCPAP vs. nHF	1.11 [0.84–1.47]	–
Methylxanthines	0.48 [0.32–0.71]	4 [2–7]
DOXAPRAM	0.80 [0.22–2.97]	–
NIPPV vs. NCPAP	0.70 [0.60–0.81]	8 [5–13]
NS-NIPPV or Bi-PAP vs. NCPAP	0.64 [0.44–0.95]	8 [4–50]
sNIPPV vs. NCPAP	0.25 [0.15–0.41]	4 [2–5]
NS-NIPPV or sNIPPV vs. NCPAP	0.28 [0.18–0.43]	4 [2–5]

NS-NIPPV, non-synchronized NIPPV; sNIPPV, synchronized NIPPV; nHF, High flow nasal cannula. Ferguson et al. (75).

The 2017 Cochrane meta-analysis compared NIPPV and NCPAP for respiratory support post-extubation and included 10 studies with 1,431 infants and reported decreased rates of respiratory failure [RR: 0.70 (95% CI 0.60, 0.80)] and reintubation rates [RR: 0.76 (95% CI 0.65, 0.88)] with NIPPV without increase in gastrointestinal side effects (59). NIPPV reduced the incidence of extubation failure and the need for re-intubation within 48 h to 1 week more effectively than NCPAP; however, it had no effect on BPD or mortality (59, 71–74). In a recent systematic review, Ferguson et al. concluded that NIPPV is superior to NCPAP in preventing extubation failure [(RR 0.70, 95% CI 0.60, 0.81; NNT 8; 95% CI 5, 13)] (**Table 4**) (75).

SYNCHRONIZED NIPPV

A clinical report by the American Academy of Pediatrics concluded that synchronized NIPPV (sNIPPV) decreases the frequency of extubation failure but the evidence for non-sNIPPV or Bi-PAP is inconclusive (76). The main reason for the absence of evidence is directly attributed to the lack of approved devices to provide an effective synchronization during NIV in USA. The most studied system for synchronization during NIPPV in newborns is the Graseby capsule (77, 78), but this system is no longer available. At present, there

are no devices in the United States that are capable of providing sNIPPV, except for neurally adjusted ventilatory assist (NAVA). However, there are devices available in other parts of the world where flow synchronization as well as Graseby capsule have been successfully used to provide sNIPPV (79).

HFNC FOR PRIMARY RESPIRATORY SUPPORT IN THE NICU

In a recent international, multicenter, randomized, non-inferiority trial, 564 preterm infants with gestational age >28 weeks (HISPTER trial) were randomized to HFNC or NCPAP. When used as primary support in preterm infants with respiratory distress, HFNC use resulted in significantly higher rates of treatment failure than NCPAP (25.5 vs. 13.3%) (80). Systematic reviews including 2016 Cochrane review and a more recent systematic review reported CPAP was superior to HFNC in preventing treatment failure and intubation [RR 1.83 (95% CI 1.43, 2.35)] in favor of CPAP (81, 82). One pilot study of 76 infants <35 weeks GA and >1,000 g birth weight compared HFNC with NIPPV as a primary mode of respiratory support and found no difference in rate of intubation and MV; however, HFNC was associated with longer duration of oxygen support (83).

HFNC VS. NCPAP IN THE NICU

In one of the largest retrospective study of 2,487 extremely preterm infants, Taha et al. reported that HFNC use was associated with higher risk of death or BPD and longer length of stay when compared to NCPAP (84). The 2016 Cochrane review of HFNC compared with NCPAP to prevent extubation failure included six trials (934 infants) and found no difference in the rate of treatment failure [RR 1.21 (95% CI 0.95, 1.55)] or reintubation RR: 0.91 (95% CI 0.68, 1.20)] within 7 days, but reported a lower rate of nasal trauma [(RR 0.64 (95% CI 0.51, 0.79)] (81). A more recent systematic review included 3 more trials and found similarly no difference in rate of treatment failure [RR 1.21 (95% CI 0.97, 1.50)] and intubation rate [RR 0.98 (95% CI 0.77, 1.24)]. However, majority of the studies included

infants greater than 28 weeks' gestation and currently there is insufficient evidence to support use of HFNC as primary mode or for post-extubation respiratory support in infants less than 28 weeks' gestation (82).

NON-INVASIVE NEURALLY ADJUSTED VENTILATORY ASSIST (NIV-NAVA)

Mechanism

NAVA is a newer mode of ventilation that utilizes electrical activity of diaphragm (Edi) using a special nasogastric tube embedded with electrodes to provide synchronized breaths (85). An electrical signal is generated in the respiratory center in the brainstem and travels via the phrenic nerve to stimulate the diaphragm. The Edi catheter with electrodes is inserted and adjusted in the esophagus to provide an optimal signal from diaphragm. Edi max as well as Edi min values are detected by the electrodes and transmitted to the ventilator. The ventilator assists the spontaneous breath by delivering a proportional pressure as determined by NAVA level. The PIP delivered is proportional to the amount of Edi. Systematically increasing NAVA levels increases PIP while maintaining Edi until the breakpoint is reached. Further increases in NAVA leads to decrease in Edi. This breakpoint is increased after extubation in premature infants (86). The initiation, duration, size, and termination of breath are controlled by the patient, and thus, potentially offering full synchronization (85). Typical settings of NIV NAVA are shown in the **Table 5**. NAVA levels are typically adjusted to keep Edi peak goal of 5–15 μ V and Edi min is kept usually between 2 and 4 μ V. If the Peak Edi is too high, NAVA level is increased to reduce the patient's work of breathing whereas if the Peak Edi is too low, the NAVA level is reduced and weaning considered. If Edi min is too high, then additional PEEP is provided whereas for low Edi min PEEP is reduced. Weaning the patient is considered after a decline in the Edi signal and peak pressure essentially showing improvement in diaphragm performance. Detailed guide for initial set up of NIV-NAVA can be accessed at <https://www.neonatologytoday.net/newsletters/nt-apr12.pdf>.

Use of NIV NAVA in NICU

Stein et al. in a retrospective study reported that in preterm infants managed on NAVA mode maintained better blood

TABLE 5 | NIV NAVA suggested settings.

	Initial settings	Maximum	Wean	Minimum
NAVA level Edi max between 5–15 μ V	2 cm H ₂ O/ μ V	4 cm H ₂ O/ μ V	0.2 to 0.5 cmH ₂ O/ μ V For Edi max > 15 μ V	0.5 cmH ₂ O/ μ V
PEEP	6 cmH ₂ O	8–10 cmH ₂ O	1 cmH ₂ O	5 cmH ₂ O
Edi Trigger	0.5 μ V	2 μ V	Adjust as needed	
Backup Pressure Control above PEEP	15 cmH ₂ O	30–35 cmH ₂ O	1–2 cmH ₂ O	Per NIPPV
Rate	40 /min	Per NIPPV	Per NIPPV	Per NIPPV
Inspiratory time	0.5 s	–	–	–
Trigger sensitivity	1 to 2	–	–	–

Edi, electrical activity of diaphragm; PEEP, positive end expiratory pressure; NIPPV, nasal intermittent positive pressure ventilation.

TABLE 6 | Studies comparing Noninvasive neutrally adjusted ventilator assist (NIV NAVA) to other forms of non-invasive ventilation.

References	Type of study	Comparison	GA (n)	Reintubation	CO ₂ clearance	Syn	Complications	Oxygen requirement/IMV duration	Outcome (Death/BPD)
Lee et al. (90)	Retrospective	NCPAP	<30 (30)	$P = 0.04$	NS	–	–	NS	NS
Kallio et al. (91)	Prospective	NCPAP	28–36 (40)	NS	NS	NS	NS	NS	NS
Yonehara et al. (92)	Retrospective	NIPPV	<30 (34)	NS	–	–	NS	–	–
Lee et al. (93)	Observational crossover	NIV-PS	<32 (15)	–	–	$P < 0.001$	–	–	–
Gibu et al. (16)	Observational crossover	NIMV	<37 (11)	–	NS	$P < 0.01$	$P < 0.01$	$P < 0.01$	–
Yagui et al. (94)	Randomized controlled	NCPAP	Preterm <1,500 g (123)	NS	–	–	NS	$P < 0.01$	NS
Yagui et al. (95)	Retrospective	NCPAP	ELBW	$P = 0.02$	–	–	NS	$P = 0.02$	NS

NIPPV, Nasal intermittent positive pressure ventilation; NCPAP, Nasal continuous positive pressure ventilation; NIV-PS, Non-invasive ventilation-Pressure support; GA, Gestational age. IMV: invasive mechanical ventilation; Syn-synchronization.

gases with lower PIP and oxygen requirements compared to synchronized IMV plus pressure support (SIMV+PS) mode of ventilation (87). Lee et al. in a randomized crossover study also reported that NAVA lowered PIP and reduced respiratory muscle load in preterm infants when compared to SIMV+PS (88). Kallio et al. performed a randomized controlled trial in 60 infants between 28 and 36 weeks GA comparing NAVA and conventional ventilation and found no difference in duration of invasive ventilation (89). Studies using NIV-NAVA mode in extremely preterm infants are limited with some of the recent studies showing promising results (Table 6) (16, 90–95). Larger trials are needed to determine if NIV-NAVA is a better mode to provide sNIPPV to prevent BPD.

NASAL HIGH FREQUENCY VENTILATION (NHFV)

To decrease the need for intubation and improve ventilation in infants with hypercarbia, using NHFV modalities such as nasal high frequency flow interrupter (NHFFI), nasal high frequency oscillatory ventilator (NHFOV) (96), nasal high frequency percussive ventilator (NHFPV) or nasal high frequency jet ventilator (NHFJV) using standard nasal interfaces have been reported. Three variables that impact the delivery of tidal volume are inspiratory time (IT), amplitude, and frequency. Longer IT, higher amplitude, and lower frequency are associated with larger tidal volume delivery during NHFV. Addition of NIPPV breaths during NHFJV also improves ventilation (97). NHFV using HFFI device, Infant Star was first reported in 1998. In this observational study of 21 preterm infants, significant improvement in ventilation was seen after starting NHFFI (98). Another study describing successful use of NHFFI in 14 patients was reported in 2008 (99).

USE OF NHFV IN NICU

In a randomized, controlled trial comparing NHFPV with NCPAP in 40 term neonates delivered by cesarean section, with a diagnosis of transient tachypnea of the newborn (TTN), NHFPV was well-tolerated and more effective in improving oxygenation when compared with NCPAP (100). Using a nasopharyngeal tube to deliver NHFOV in 20 preterm neonates during weaning from IMV, NHFOV was successfully used in 91% of the patients at first attempt at extubation (101). Mukerji et al. reviewed 52 patients treated with rescue NHFOV when other NIV modes failed. Intubation was avoided in 58% of the cases (102). In a small randomized, controlled trial involving 39 patients with a birth weight <1,250 g, NHFOV was found to be not superior to Si-PAP (103). Most likely reason for lack of success in this pilot study was the use of lower MAP in the NHFOV group. In a recent meta-analysis of 8 RCTs involving 463 patients, NHFOV significantly improved CO₂ clearance and reduced the need for intubation compared with NCPAP/bi-phasic CPAP (104). There are no clinical trials using NHFJV. There is only one reported case series showing successful use of NHFJV in selected extremely preterm infants immediately after extubation from IMV (105). In a review of 6 NHFV studies involving 111 patients; different inspiratory time, amplitude, and frequency were used (96). Suggested settings for NHFV are shown in Table 7.

COMPLICATIONS OF USING NIV

Even if NIV offers a number of benefits over IMV some extremely preterm infants may develop complications while receiving NIV. Majority of complications are related to injury to nasal mucosa as well as nasal septum.

TABLE 7 | Suggested settings for Nasal High Frequency Ventilation.

Frequency, Hz	Start at 6–8 Hz; May decrease to 4 Hz in patients with hypercapnia; If using HFJV, start at 300 bpm (5 Hz) and may decrease to 240 bpm (4 Hz)
Amplitude, cmH ₂ O	MAPx2; Start at 20–30 cmH ₂ O; May increase to as high 70 cmH ₂ O. If using during weaning, set Amplitude equaling PIP prior to extubation
I: E ratio	Start at 1:1; May change to 1:2 in cases of gas trapping; If using HFJV, jet valve on time: 20 ms and may increase to 30–34 ms to improve oxygenation and increase tidal volume delivery
Mean Airway Pressure (MAP), cmH ₂ O	MAP: Start with the same MAP as on SIMV or 2–3 cmH ₂ O higher than CPAP; Start at 8–10 cmH ₂ O; May increase as needed based on FiO ₂ and or lung expansion
NIPPV Back up rate	If available, use rates between 30 and 40 bpm; If using HFJV, keep the NIPPV settings same as before adding NHFJV

I:E ratio, Inspiration: Expiration ratio; HFJV, high frequency jet ventilator; NIPPV, nasal intermittent positive pressure ventilation; MAP, Mean Airway Pressure; SIMV, Synchronized intermittent mandatory ventilation.

NASAL INJURIES DURING NIV

One of the major problems with NIV use in the NICU in the occurrence of septal or nasal mucosal injuries resulting in nasal deformities. Snugly fit nasal prongs may put extremely preterm infants at risk for causing nasal trauma including erythema or blanching, ulceration, and columellar necrosis. Reported incidence of these complications varies from 20 to 60% in neonates. Both frequency and severity of nasal trauma has been shown to be higher in infants at lower gestational age (>90% in neonates <28 weeks of gestational age), lower birth weight, longer duration of NCPAP and longer NICU stay (106). It is important to choose the right interface with correct size of prongs as well as fittings as onset of nasal injury to the columella has been reported to occur within a mean of 2–3 days of CPAP commencement, and in some cases occurring as early as 18 h after commencement. The use of nasal barrier dressings and nasal masks as an alternative to binasal prongs may be effective interventions to reduce nasal injury. HFNC causes less nasal injury than CPAP, but it may not provide sufficient respiratory support for the smallest, sickest preterm infants (107). These complications not only have cosmetic or functional sequelae but also place the infants at risk for developing nosocomial infections. In one study nasal breakdown with the INCA prongs and subsequent use of the RAM NC did not worsen or contribute nasal injuries. Also, there were no new instances of nasal breakdown or injury reported with use of the RAM NC (108). A recent study showed skin or mucosal breakdown with RAM NC was significantly lower compared to other nasal interfaces (8 vs. 53%, $P < 0.001$) (109). Common recommendations for prevention of nasal trauma due to NCPAP in neonates include careful monitoring of the nose, avoidance of pressure, friction, and moisture.

OTHER COMPLICATIONS WITH NIV

Systemic complications related to NIV use are usually rare and account for less than 5% of patients. Pneumothorax can

occur in acute phase and it is most commonly related to underlying lung disease rather than NIV itself. Also, after surfactant administration with sudden change in the compliance may lead to air leaks as well. HFNC may have reduced occurrence of pneumothorax compared to CPAP. Small pneumothoraces usually resolves spontaneously, and one may not need to change the modality if infant is otherwise stable. Rarely an intervention is needed to evacuate the air to re-inflate the affected lung. A recent study showed a decrease risk of pneumothoraces after implementing NIV in the delivery room instead of intubation in extremely low birth weight infants (3).

It is common to observe abdominal distension with or without feeding intolerance with NIV. Some infants may need to have orogastric tube to vent the stomach and to evacuate air. There may be transient feeding intolerance. Decrease in flow may help relieve gastric distension as well. Infants who have undergone upper gastrointestinal surgery are at higher risk of complications, such as leak at the site of anastomosis if NIV is used in the immediate post-operative period (110). In a small case series, Pandita et al. reported facial palsy in 3 patients, who were on NCPAP and speculated that pressure over the stylomastoid foramen by the NCPAP interface might have contributed to ipsilateral facial palsy (111).

CONCLUSIONS

In extremely preterm infants, optimal pulmonary outcomes could be achieved by minimizing the duration of IMV. NIV is currently best provided by early use of NIPPV from DR through 32 to 33 weeks postmenstrual age in the NICU. NCPAP may be used when weaning from NIPPV, followed by low flow nasal cannula (<2 LPM) in extremely preterm infants to minimize lung injury. sNIPPV could be delivered using NIV NAVA but needs more evidence to support its use in this specific population. NHFV using NHFOV as well as NHFJV have the potential as a rescue mode for use in this population especially when the lung disease is severe requiring higher pressure (PIP >30 cmH₂O) to improve gas exchange.

AUTHOR CONTRIBUTIONS

YS: Drafting of the initial manuscript, critical revision of the manuscript for important intellectual content. RR: Review of the literature, creation of tables, checking and adding references, creation of figures, drafting and subsequent revisions of the manuscript for important intellectual content. MB: Review of the literature, creation of tables, checking and adding references, drafting and subsequent revisions of the manuscript for important intellectual content. HM: Review of the literature, creation of tables, drafting and subsequent revisions of the manuscript for important intellectual content. All Authors reviewed the final version of this manuscript and approved for submission.

REFERENCES

- Walsh MC, Morris BH, Wrage LA, Vohr BR, Poole WK, Tyson JE, et al. Extremely low birth weight neonates with protracted ventilation: mortality and 18-month neurodevelopmental outcomes. *J Pediatr.* (2005) 146:798–804. doi: 10.1016/j.jpeds.2005.01.047
- Zhang H, Dysart K, Kendrick DE, Li L, Das A, Hintz SR, et al. Prolonged respiratory support of any type impacts outcomes of extremely low birth weight infants. *Pediatr Pulmonol.* (2018) 53:1447–55. doi: 10.1002/ppul.24124
- Govindaswami B, Nudelman M, Narasimhan SR, Huang A, Misra S, Urquidez G, et al. Eliminating risk of intubation in very preterm infants with noninvasive cardiorespiratory support in the delivery room and neonatal intensive care unit. *Biomed Res Int.* (2019) 2019:5984305. doi: 10.1155/2019/5984305
- Stoll BJ, Hansen NI, Bell EF, Walsh MC, Carlo WA, Shankaran S, et al. Trends in care practices, morbidity, and mortality of extremely preterm neonates, 1993–2012. *JAMA.* (2015) 314:1039–51. doi: 10.1001/jama.2015.10244
- Laptook AR, O'Shea TM, Shankaran S, Bhaskar B, NICHD Neonatal Network. Adverse neurodevelopmental outcomes among extremely low birth weight infants with a normal head ultrasound: prevalence and antecedents. *Pediatrics.* (2005). 115:673–80. doi: 10.1542/peds.2004-0667
- Cheong JLY, Doyle LW. An update on pulmonary and neurodevelopmental outcomes of bronchopulmonary dysplasia. *Semin Perinatol.* (2018) 42:478–84. doi: 10.1053/j.semperi.2018.09.013
- Jobe AH, Bancalari E. Bronchopulmonary dysplasia. *Am J Respir Crit Care Med.* (2001) 163:1723–9. doi: 10.1164/ajrccm.163.7.2011060
- 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) 79:26–30.
- Jensen EA, Dysart K, Gantz MG, McDonald S, Bamat NA, Keszler M, et al. The diagnosis of bronchopulmonary dysplasia in very preterm infants. An evidence-based approach. *Am J Respir Crit Care Med.* (2019) 200:751–9. doi: 10.1164/rccm.201812-2348OC
- Verder H, Robertson B, Greisen G, Ebbesen F, Albertsen P, Lundstrøm K, et al. Surfactant therapy and nasal continuous positive airway pressure for newborns with respiratory distress syndrome. Danish-Swedish Multicenter Study Group. *N Engl J Med.* (1994) 331:1051–5. doi: 10.1056/NEJM199410203311603
- Jena SR, Bains HS, Pandita A, Verma A, Gupta V, Kallem VR, et al. Surfactant therapy in premature babies: SurE or InSurE. *Pediatr Pulmonol.* (2019) 54:1747–52. doi: 10.1002/ppul.24479
- Pandita A, Panza R. Sure or insure. *Pediatr Pulmonol.* (2020) 55:18–9. doi: 10.1002/ppul.24573
- Kribs A, Roll C, Göpel W, Wieg C, Groneck P, Laux R, et al. Nonintubated surfactant application vs conventional therapy in extremely preterm infants: a randomized clinical trial. *JAMA Pediatr.* (2015) 169:723–30. doi: 10.1001/jamapediatrics.2015.0504
- De Paoli AG, Morley C, Davis PG. Nasal CPAP for neonates: what do we know in 2003? *Arch Dis Child Fetal Neonatal Ed.* (2003) 88:F168–72. doi: 10.1136/fn.88.3.F168
- Kiciman NM, Andréasson B, Bernstein G, Mannino FL, Rich W, Henderson C, et al. Thoracoabdominal motion in newborns during ventilation delivered by endotracheal tube or nasal prongs. *Pediatr Pulmonol.* (1998) 25:175–81. doi: 10.1002/(SICI)1099-0496(199803)25:3<175::AID-PPUL7>3.0.CO;2-L
- Gibu CK, Cheng PY, Ward RJ, Castro B, Heldt GP. Feasibility and physiological effects of noninvasive neurally adjusted ventilatory assist in preterm infants. *Pediatr Res.* (2017) 82:650–7. doi: 10.1038/pr.2017.100
- Frizzola M, Miller TL, Rodriguez ME, Zhu Y, Rojas J, Heseck A, et al. High-flow nasal cannula: impact on oxygenation and ventilation in an acute lung injury model. *Pediatr Pulmonol.* (2011) 46:67–74. doi: 10.1002/ppul.21326
- Stefanescu BM, Murphy W, Hansell BJ, Fuloria M, Morgan TM, Aschner JL. A randomized, controlled trial comparing two different continuous positive airway pressure systems for the successful extubation of extremely low birth weight infants. *Pediatrics.* (2003) 112:1031–8. doi: 10.1542/peds.112.5.1031
- Gupta S, Sinha SK, Tin W, Donn SM. A randomized controlled trial of post-extubation bubble continuous positive airway pressure versus infant flow driver continuous positive airway pressure in preterm infants with respiratory distress syndrome. *J Pediatr.* (2009) 154:645–50. doi: 10.1016/j.jpeds.2008.12.034
- Weydig H, Ali N, Kakkilaya V. Noninvasive ventilation in the delivery room for the preterm infant. *Neoreviews.* (2019) 20:e489–99. doi: 10.1542/neo.20-9-e489
- Finer NN, Rich W, Wang C, Leone T. Airway obstruction during mask ventilation of very low birth weight infants during neonatal resuscitation. *Pediatrics.* (2009) 123:865–69. doi: 10.1542/peds.2008-0560
- Schmolzer GM, Dawson JA, Kamlin CO, O'Donnell CP, Morley CJ, Davis PG. Airway obstruction and gas leak during mask ventilation of preterm infants in the delivery room. *Arch Dis Child Fetal Neonatal Ed.* (2011) 96:F254–57. doi: 10.1136/adc.2010.191171
- van Vonderen JJ, Kleijn TA, Schilleman K, Walther FJ, Hooper SB, te Pas AB. Compressive force applied to a manikin's head during mask ventilation. *Arch Dis Child Fetal Neonatal Ed.* (2012) 97:F254–8. doi: 10.1136/archdischild-2011-300336
- Murthy V, Dattani N, Peacock JL, Fox GF, Campbell ME, Milner AD, et al. The first five inflations during resuscitation of prematurely born infants. *Arch Dis Child Fetal Neonatal Ed.* (2012) 97:F249–53. doi: 10.1136/archdischild-2011-300117
- Kilmartin KC, Finn D, Hawkes GA, Kenosi M, Dempsey EM, Livingstone V, et al. Corrective ventilation strategies in delivery room resuscitation of preterm infants. *Acta Paediatr.* (2018) 107:2066–70. doi: 10.1111/apa.14570
- Yang KC, Te Pas AB, Weinberg DD, Foglia EE. Corrective steps to enhance ventilation in the delivery room. *Arch Dis Child Fetal Neonatal Ed.* (2020) 2019:318579. doi: 10.1136/archdischild-2019-318579
- Capasso L, Capasso A, Raimondi F, Vendemmia M, Araimo G, Paludetto R. A randomized trial comparing oxygen delivery on intermittent positive pressure with nasal cannulae versus facial mask in neonatal primary resuscitation. *Acta Paediatr.* (2005) 94:197–200. doi: 10.1080/08035250410025113
- Biniwale M, Wertheimer F. Decrease in delivery room intubation rates after use of nasal intermittent positive pressure ventilation in the delivery room for resuscitation of very low birth weight infants. *Resuscitation.* (2017) 116:33–8. doi: 10.1016/j.resuscitation.2017.05.004
- Petrillo F, Valenzano L, Franco C, Calò G, Denticò D, Manzoni P, et al. Pulmonary recruitment strategy in preterm neonates < 29 weeks of gestational age to reduce the need for intubation in the delivery room. *Am J Perinatol.* (2019) 36:S115–9. doi: 10.1055/s-0039-1692134
- Ramanathan R, Biniwale M. Noninvasive ventilation and high-flow nasal cannula. In: Rajeev PK, Lakshminrusimha S, Vidyasagar D, editors. *Essentials of Neonatal Ventilation 1st Edition*. New Delhi: Elsevier (2018). p. 250–79.
- Wyckoff MH, Aziz K, Escobedo MB, Kapadia VS, Kattwinkel J, Perlman JM, et al. Part 13: neonatal resuscitation: 2015 American heart association guidelines update for cardiopulmonary resuscitation and emergency cardiovascular care. *Circulation.* (2015) 132:S543–60. doi: 10.1161/CIR.0000000000000267
- Green EA, Dawson JA, Davis PG, De Paoli AG, Roberts CT. Assessment of resistance of nasal continuous positive airway pressure interfaces. *Arch Dis Child Fetal Neonatal Ed.* (2019) 104:F535–9. doi: 10.1136/archdischild-2018-315838
- Jasani B, Ismail A, Rao S, Patole S. Effectiveness and safety of nasal mask versus binasal prongs for providing continuous positive airway pressure in preterm infants—A systematic review and meta-analysis. *Pediatr Pulmonol.* (2018) 53:987–92. doi: 10.1002/ppul.24014
- Iyer NP, Chatburn R. Evaluation of a nasal cannula in noninvasive ventilation using a lung simulator. *Respir Care.* (2015) 60:508–12. doi: 10.4187/respcare.03560
- Roberts CT, Kortekaas R, Dawson JA, Manley BJ, Owen LS, Davis PG. The effects of non-invasive respiratory support on oropharyngeal temperature and humidity: a neonatal manikin study. *Arch Dis Child Fetal Neonatal Ed.* (2016) 101:F248–52. doi: 10.1136/archdischild-2015-308991
- Ullrich TL, Czernik C, Bühner C, Schmalisch G, Fischer HS. Nasal high-frequency oscillatory ventilation impairs heated humidification: a neonatal bench study. *Pediatr Pulmonol.* (2017) 52:1455–60. doi: 10.1002/ppul.23824

37. Heldt GP, McLroy MB. Distortion of chest wall and work of diaphragm in preterm infants. *J Appl Physiol.* (1987) 62:164–9. doi: 10.1152/jappl.1987.62.1.164
38. 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) 114:651–7. doi: 10.1542/peds.2004-0394
39. Morley CJ, Davis PG, Doyle LW, Brion LP, Hascoet JM, Carlin JB, et al. Nasal CPAP or intubation at birth for very preterm infants. *N Engl J Med.* (2008) 358:700–8. doi: 10.1056/NEJMoa072788
40. SUPPORT Study Group of the Eunice Kennedy Shriver NICHD Neonatal Research Network, Finer NN, Carlo WA, Walsh MC, Rich W, Gantz MG, et al. Early CPAP versus surfactant in extremely preterm infants. *N Engl J Med.* (2010) 362:1970–9. doi: 10.1056/NEJMoa0911783
41. Dunn MS, Kaempf J, de Klerk A, de Klerk R, Reilly M, Howard D, et al. Randomized trial comparing 3 approaches to the initial respiratory management of preterm neonates. *Pediatrics.* (2011) 128:e1069–76. doi: 10.1542/peds.2010-3848
42. Rojas MA, Lozano JM, Rojas MX, Laughon M, Bose CL, Rondon MA, et al. Very early surfactant without mandatory ventilation in premature infants treated with early continuous positive airway pressure: a randomized, controlled trial. *Pediatrics.* (2009) 123:137–42. doi: 10.1542/peds.2007-3501
43. Sandri F, Plavka R, Ancora G, Simeoni U, Stranak Z, Martinelli S et al. Prophylactic or early selective surfactant combined with nCPAP in very preterm infants. *Pediatrics.* (2010) 125:e1402–9. doi: 10.1542/peds.2009-2131
44. Göpel W, Kribs A, Ziegler A, Laux R, Hoehn T, Wieg C, et al. Avoidance of mechanical ventilation by surfactant treatment of spontaneously breathing preterm infants (AMV): an open-label, randomised, controlled trial. *Lancet.* (2011) 378:1627–34. doi: 10.1016/S0140-6736(11)60986-0
45. Kanmaz HG, Erdeve O, Canpolat FE, Mutlu B, Dilmen U. Surfactant administration via thin catheter during spontaneous breathing: randomized controlled trial. *Pediatrics.* (2013) 131:e502–9. doi: 10.1542/peds.2012-0603
46. Tapia JL, Urzua S, Bancalari A, Meritano J, Torres G, Fabres J, et al. Randomized trial of early bubble continuous positive airway pressure for very low birth weight infants. *J Pediatr.* (2012) 161:75–80. doi: 10.1016/j.jpeds.2011.12.054
47. Fischer HS, Bührer C. Avoiding endotracheal ventilation to prevent bronchopulmonary dysplasia: A Meta-analysis. *Pediatrics.* (2013) 132:e1351–61. doi: 10.1542/peds.2013-1880
48. Schmölzer GM, Kumar M, Pichler G, Aziz K, O'Reilly M, Cheung PY. Non-invasive versus invasive respiratory support in preterm infants at birth: systematic review and meta-analysis. *BMJ.* (2013) 347:f5980. doi: 10.1136/bmj.f5980
49. Doyle LW, Carse E, Adams AM, Ranganathan S, Opie G, Cheong JLY. Ventilation in extremely preterm infants and respiratory function at 8 Years. *N Engl J Med.* (2017). 377:329–37. doi: 10.1056/NEJMoa1700827
50. te Pas AB, Siew M, Wallace MJ, Kitchen MJ, Fouras A, Lewis RA, et al. Establishing functional residual capacity at birth: the effect of sustained inflation and positive end-expiratory pressure in a preterm rabbit model. *Pediatr Res.* (2009) 65:537–41. doi: 10.1203/PDR.0b013e31819da21b
51. Kirpalani H, Ratcliffe SJ, Kesler M, Davis PG, Foglia EE, Te Pas A, et al. Effect of Sustained Inflation vs Intermittent Positive Pressure Ventilation on Bronchopulmonary Dysplasia or Death among Extremely Preterm Infants: The Sail Randomized Clinical Trial. *JAMA.* (2019) 321:1165–75. doi: 10.1001/jama.2019.1660
52. Bruschetti M, O'Donnell CP, Davis PG, Morley CJ, Moja L, Calevo MG. Sustained versus standard inflations during neonatal resuscitation to prevent mortality and improve respiratory outcomes. *Cochrane Database Syst Rev.* (2020) 3:CD004953. doi: 10.1002/14651858.CD004953.pub4
53. Foglia EE, Te Pas AB, Kirpalani H, Davis PG, Owen LS, van Kaam AH, et al. Sustained inflation vs standard resuscitation for preterm infants: a systematic review and meta-analysis. *JAMA Pediatr.* (2020) 3:e195897. doi: 10.1001/jamapediatrics.2019.5897
54. Reynolds P, Leontiadi S, Lawson T, Otunla T, Ejiwumi O, Holland N. Stabilisation of premature infants in the delivery room with nasal high flow. *Arch Dis Child Fetal Neonatal Ed.* (2016) 101:F284–7. doi: 10.1136/archdischild-2015-309442
55. Rojas-Reyes MX, Morley CJ, Soll R. Prophylactic versus selective use of surfactant in preventing morbidity and mortality in preterm infants. *Cochrane Database Syst Rev.* (2012) 3:CD000510. doi: 10.1002/14651858.CD000510.pub2
56. 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) 147:341–7. doi: 10.1016/j.jpeds.2005.04.062
57. Dargaville PA, Gerber A, Johansson S, De Paoli AG, Kamlin CO, Orsini F, et al. Incidence and outcome of CPAP failure in preterm infants. *Pediatrics.* (2016) 138:e20153985. doi: 10.1542/peds.2015-3985
58. Lemyre B, Davis PG, de Paoli AG. Nasal intermittent positive pressure ventilation (NIPPV) versus nasal continuous positive airway pressure (NCPAP) for apnea of prematurity. *Cochrane Database Syst Rev.* (2002) 1:CD002272. doi: 10.1002/14651858.CD002272
59. Lemyre B, Davis PG, De Paoli AG, Kirpalani H. Nasal intermittent positive pressure ventilation (NIPPV) versus nasal continuous positive airway pressure (NCPAP) for preterm neonates after extubation. *Cochrane Database of Syst Rev.* (2017) 2:CD003212. doi: 10.1002/14651858.CD003212.pub3
60. Armanian AM, Badiee Z, Heidari G, Feizi A, Salehimehr N. Initial treatment of respiratory distress syndrome with nasal intermittent mandatory ventilation versus nasal continuous positive airway pressure: a randomized controlled trial. *Int J Prev Med.* (2014) 5:1543–51.
61. Bisceglia M, Belcastro A, Poerio V, Raimondi F, Mesuraca L, Crugliano C, et al. A comparison of nasal intermittent versus continuous positive pressure delivery for the treatment of moderate respiratory distress syndrome in preterm infants. *Minerva Pediatr.* (2007) 59:91–5.
62. Kirpalani H, Millar D, Lemyre B, Yoder BA, Chiu A, Roberts RS, et al. A trial comparing noninvasive ventilation strategies in preterm infants. *N Engl J Med.* (2013) 369:611–20. doi: 10.1056/NEJMoa1214533
63. Meneses J, Bhandari V, Alves JG, Herrmann D. Noninvasive ventilation for respiratory distress syndrome: a randomized controlled trial. *Pediatrics.* (2011) 127:300–7. doi: 10.1542/peds.2010-0922
64. Kugelmann A, Feferkorn I, Riskin A, Chistyakov I, Kaufman B, Bader D. Nasal intermittent mandatory ventilation versus nasal continuous positive airway pressure for respiratory distress syndrome: a randomized, controlled, prospective study. *J Pediatr.* (2007) 150:521–6. doi: 10.1016/j.jpeds.2007.01.032
65. Sai Sunil Kishore M, Dutta S, Kumar P. Early nasal intermittent positive pressure ventilation versus continuous positive airway pressure for respiratory distress syndrome. *Acta Paediatr.* (2009) 98:1412–5. doi: 10.1111/j.1651-2227.2009.01348.x
66. Salama GS, Ayyash FF, Al-Rabadi AJ, Alquran ML, Shakkoury AG. Nasal-IMV versus nasal-CPAP as an initial mode of respiratory support for premature infants with RDS: a prospective randomized clinical trial. *Rawal Med J.* (2015) 40:197–202.
67. Lista G, Castoldi F, Fontana P, Daniele I, Cavigioli F, Rossi S, et al. Nasal continuous positive airway pressure (CPAP) versus bi-level nasal CPAP in preterm babies with respiratory distress syndrome: a randomised control trial. *Arch Dis Child Fetal Neonatal Ed.* (2010) 95:F85–9. doi: 10.1136/adc.2009.169219
68. Ramanathan R, Sekar KC, Rasmussen M, Bhatia J, Soll RF. Nasal intermittent positive pressure ventilation after surfactant treatment for respiratory distress syndrome in preterm infants <30 weeks' gestation: a randomized, controlled trial. *J Perinatol.* (2012) 32:336–43. doi: 10.1038/jp.2012.1
69. Oncel MY, Arayici S, Uras N, Alyamac-Dizdar E, Sari FN, Karahan S, et al. Nasal continuous positive airway pressure versus nasal intermittent positive-pressure ventilation within the minimally invasive surfactant therapy approach in preterm infants: a randomized controlled trial. *Arch Dis Child Fetal Neonatal Ed.* (2016) 101:F423–8. doi: 10.1136/archdischild-2015-308204
70. Lemyre B, Laughon M, Bose C, Davis PG. Early nasal intermittent positive pressure ventilation (NIPPV) versus early nasal continuous positive airway pressure (NCPAP) for preterm infants. *Cochrane Database Syst Rev.* (2016) 12:CD005384. doi: 10.1002/14651858.CD005384.pub2
71. Friedlich P, Lecart C, Posen R, Ramicone E, Chan L, Ramanathan R. A randomized trial of nasopharyngeal-synchronised intermittent mandatory ventilation versus nasopharyngeal continuous positive airway pressure in

- very low birth weight infants following extubation. *J Perinatol.* (1999) 19:413–8. doi: 10.1038/sj.jp.7200205
72. Gao WW, Tan SZ, Chen YB, Zhang Y, Wang Y. Randomized trial of nasal synchronized intermittent mandatory ventilation compared with nasal continuous positive airway pressure in preterm infants with respiratory distress syndrome. *Zhongguo Dang Dai Er Ke Za Zhi.* (2010) 12:524–6.
 73. Jasani B, Nanavati R, Kabra N, Rajdeo S, Bhandari V. Comparison of non-synchronized nasal intermittent positive pressure ventilation versus nasal continuous positive airway pressure as post-extubation respiratory support in preterm infants with respiratory distress syndrome: a randomized controlled trial. *J Matern Fetal Neonatal Med.* (2016) 29:1546–51. doi: 10.3109/14767058.2015.1059809
 74. Kahramaner Z, Erdemir A, Turkoglu E, Cosar H, Sutcuoglu S, Ozer EA. Unsynchronized nasal intermittent positive pressure versus nasal continuous positive airway pressure in preterm infants after extubation. *J Matern Fetal Neonatal Med.* (2014) 27:926–9. doi: 10.3109/14767058.2013.846316
 75. Ferguson KN, Roberts CT, Manley BJ, Davis PG. Interventions to improve rates of successful extubation in preterm infants. A systematic review and meta-analysis. *JAMA Pediatr.* (2017) 171:165–74. doi: 10.1001/jamapediatrics.2016.3015
 76. Cummings JJ, Polin R. Committee on fetus and newborn, american academy of pediatrics. Noninvasive respiratory support. *Pediatrics.* (2016) 137:e20153758. doi: 10.1542/peds.2015-3758
 77. Barrington KJ, Finer NN, Bull D. Randomised controlled trial of nasal synchronized intermittent mandatory ventilation compared with continuous positive airway pressure after extubation of very low birth weight infants. *Pediatrics.* (2001) 107:638–41. doi: 10.1542/peds.107.4.638
 78. Khalaf MN, Brodsky N, Hurley J, Bhandari V. A prospective randomized, controlled trial comparing synchronized nasal intermittent positive pressure ventilation versus nasal continuous positive airway pressure as modes of extubation. *Pediatrics.* (2001) 108:13–7. doi: 10.1542/peds.108.1.13
 79. Ramos-Navarro C, Sanchez-Luna M, Sanz-López E, Maderuelo-Rodriguez E, Zamora-Flores E. Effectiveness of synchronized noninvasive ventilation to prevent intubation in preterm infants. *AJP Rep.* (2016) 6:e264–71. doi: 10.1055/s-0036-1586205
 80. Roberts CT, Owen LS, Manley BJ, Froisland DH, Donath SM, Dalziel KM, et al. Nasal high-flow therapy for primary respiratory support in preterm infants. *N Engl J Med.* (2016) 375:1142–52. doi: 10.1056/NEJMoa1603694
 81. Wilkinson D, Andersen C, O'Donnell CPF, De Paoli AG, Manley BJ. High flow nasal cannula for respiratory support in preterm infants. *Cochrane Database of Syst Rev.* (2016) 2:CD006405. doi: 10.1002/14651858.CD006405.pub3
 82. Hodgson KA, Manley BJ, Davis PG. Is nasal high flow inferior to continuous positive airway pressure for neonates? *Clin Perinatol.* (2019) 46:537–51. doi: 10.1016/j.clp.2019.05.005
 83. Kugelman A, Riskin A, Said W, Shoris I, Mor F, Bader D. A randomized pilot study comparing heated humidified high-flow nasal cannulae with NIPPV for RDS. *Pediatr Pulmonol.* (2015) 50:576–83. doi: 10.1002/ppul.23022
 84. Taha DK, Kornhauser M, Greenspan JS, Dysart KC, Aghai ZH. High flow nasal cannula use is associated with increased morbidity and length of hospitalization in extremely low birth weight infants. *J Pediatr.* (2016) 173:50–5. doi: 10.1016/j.jpeds.2016.02.051
 85. Sinderby C, Beck J. Neurally adjusted ventilatory assist (NAVA): an update and summary of experiences. *Neth J Crit Care.* (2007) 11:243–52.
 86. LoVerde B, Firestone KS, Stein HM. Comparing changing neurally adjusted ventilatory assist (NAVA) levels in intubated and recently extubated neonates. *J Perinatol.* (2016) 36:1097–100. doi: 10.1038/jp.2016.152
 87. Stein H, Howard D. Neurally adjusted ventilatory assist in neonates weighing <1500 grams: a retrospective analysis. *J Pediatr.* (2012) 160:786–9. doi: 10.1016/j.jpeds.2011.10.014
 88. Lee J, Kim H, Sohn JA, Lee JA, Choi CW, Kim EK, et al. Randomized crossover study of neurally adjusted ventilatory assist in preterm infants. *J Pediatr.* (2012) 161:808–13. doi: 10.1016/j.jpeds.2012.04.040
 89. Kallio M, Koskela U, Peltoniemi O, Kontiokari T, Pokka T, Suo-Palosaari M, et al. Neurally adjusted ventilatory assist (NAVA) in preterm newborn infants with respiratory distress syndrome—a randomized controlled trial. *Eur J Pediatr.* (2016) 175:1175–83. doi: 10.1007/s00431-016-2758-y
 90. Lee BK, Shin SH, Jung YH, Kim EK, Kim HS. Comparison of NIV-NAVA and NCPAP in facilitating extubation for very preterm infants. *BMC Pediatr.* (2019) 19:298. doi: 10.1186/s12887-019-1683-4
 91. Kallio M, Mahlman M, Koskela U, Aikio O, Suo-Palosaari M, Pokka T, et al. NIV NAVA versus nasal CPAP in premature infants: a randomized clinical trial. *Neonatology.* (2019) 116:380–4. doi: 10.1159/000502341
 92. Yonehara K, Ogawa R, Kamei Y, Oda A, Kokubo M, Hiroma T, et al. Non-invasive neurally adjusted ventilatory assist versus nasal intermittent positive-pressure ventilation in preterm infants born before 30 weeks' gestation. *Pediatr Int.* (2018) 60:957–61. doi: 10.1111/ped.13680
 93. Lee J, Kim HS, Jung YH, Shin SH, Choi CW, Kim EK, et al. Non-invasive neurally adjusted ventilatory assist in preterm infants: a randomised phase II crossover trial. *Arch Dis Child Fetal Neonatal Ed.* (2015) 100:F507–13. doi: 10.1136/archdischild-2014-308057
 94. Yagui AC, Meneses J, Zólio BA, Brito GMG, da Silva RJ, Rebello CM. Nasal continuous positive airway pressure (NCPAP) or noninvasive neurally adjusted ventilatory assist (NIV-NAVA) for preterm infants with respiratory distress after birth: a randomized controlled trial. *Pediatr Pulmonol.* (2019) 54:1704–11. doi: 10.1002/ppul.24466
 95. Yagui ACZ, Gonçalves PA, Murakami SH, Santos AZ, Zacharias RSB, Rebello CM. Is noninvasive neurally adjusted ventilatory assistance (NIV-NAVA) an alternative to NCPAP in preventing extubation failure in preterm infants? *J Matern Fetal Neonatal Med.* (2019) 24:1–151. doi: 10.1080/14767058.2019.1697669
 96. De Luca D, Dell'Orto V. Non-invasive high-frequency oscillatory ventilation in neonates: review of physiology, biology and clinical data. *Arch Dis Child Fetal Neonatal Ed.* (2016) 101:F565–70. doi: 10.1136/archdischild-2016-310664
 97. Yoder BA, Albertine KH, Null DM Jr. High-frequency ventilation for non-invasive respiratory support of neonates. *Semin Fetal Neonatal Med.* (2016) 21:162–73. doi: 10.1016/j.siny.2016.02.001
 98. van der Hoeven M, Brouwer E, Blanco CE. Nasal high frequency ventilation in neonates with moderate respiratory insufficiency. *Arch Dis Child Fetal Neonatal Ed.* (1998) 79:F61–3. doi: 10.1136/fn.79.1.F61
 99. Colaizy TT, Younis UM, Bell EF, Klein JM. Nasal high-frequency ventilation for premature infants. *Acta Paediatr.* (2008) 97:1518–22. doi: 10.1111/j.1651-2227.2008.00900.x
 100. Dumas De La Roque E, Bertrand C, Tandonnet O, Rebola M, Roquand E, Renesme L, et al. Nasal high frequency percussive ventilation versus nasal continuous positive airway pressure in transient tachypnea of the newborn: a pilot randomized controlled trial (NCT00556738). *Pediatr Pulmonol.* (2011) 46:218–23. doi: 10.1002/ppul.21354
 101. Czernik C, Schmalisch G, Bühner C, Proquitté H. Weaning of neonates from mechanical ventilation by use of nasopharyngeal high-frequency oscillatory ventilation: a preliminary study. *J Matern Fetal Neonatal Med.* (2012) 25:374–8. doi: 10.3109/14767058.2011.580401
 102. Mukerji A, Singh B, Helou SE, Fusch C, Dunn M, Belik J, et al. Use of noninvasive high-frequency ventilation in the neonatal intensive care unit: a retrospective review. *Am J Perinatol.* (2015) 30:171–6. doi: 10.1055/s-0034-1381317
 103. Mukerji A, Sarmiento K, Lee B, Hassall K, Shah V. Non-invasive high-frequency ventilation versus bi-phasic continuous positive airway pressure (BP-CPAP) following CPAP failure in infants <1250 g: a pilot randomized controlled trial. *J Perinatol.* (2017) 37:49–53. doi: 10.1038/jp.2016.172
 104. Li J, Li X, Huang X, Zhang Z. Noninvasive high-frequency oscillatory ventilation as respiratory support in preterm infants: a meta-analysis of randomized controlled trials. *Respir Res.* (2019) 20:58. doi: 10.1186/s12931-019-1023-0
 105. De Beritto T, Keel JJ, Ramanathan R, Biniwale M, Cayabyab R. Nasal high frequency jet ventilation versus nasal intermittent positive pressure ventilation as a mean of post extubation respiratory support. *J Investig Med.* (2020) 68:A53.
 106. Fischer C, Bertelle V, Hohlfeld J, Forcada-Guex M, Stadelmann-Diaw C, Tolsa JF. Nasal trauma due to continuous positive airway pressure in neonates. *Arch Dis Child Fetal Neonatal Ed.* (2010) 95:F447–51. doi: 10.1136/adc.2009.179416

107. Imbulana DI, Manley BJ, Dawson JA, Davis PG, Owen LS. Nasal injury in preterm infants receiving non-invasive respiratory support: a systematic review. *Arch Dis Child Fetal Neonatal Ed.* (2018) 103:F29–35. doi: 10.1136/archdischild-2017-313418
108. Nzegwu NI, Mack T, DellaVentura R, Dunphy L, Koval N, Levit O, et al. Systematic use of the ram nasal cannula in the yale-new haven children's hospital neonatal intensive care unit: a quality improvement project. *J Matern Fetal Neonatal Med.* (2015) 28:718–21. doi: 10.3109/14767058.2014.929659
109. Drescher GS, Hughes CW. Comparison of interfaces for the delivery of noninvasive respiratory support to low birthweight infants. *Respir Care.* (2018) 10:1197–206. doi: 10.4187/respcare.05978
110. Ferrand A, Roy SK, Faure C, Moussa A, Aspirot A. Postoperative noninvasive ventilation and complications in esophageal atresia-tracheoesophageal fistula. *J Pediatr Surg.* (2019) 54:945–8. doi: 10.1016/j.jpedsurg.2019.01.023
111. Pandita A, Mishra N, Gupta G, Shukla A. Neonatal facial palsy, a case series: is CPAP the culprit? *BMJ*

Case Rep. (2018) 2018:bcr2018224842. doi: 10.1136/bcr-2018-224842

Conflict of Interest: RR has a joint patent on RAM Nasal Cannula and receives royalty payments.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2020 Shi, Muniraman, Biniwale and Ramanathan. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.