# Guidelines and Best Practices for Mask-Free NIV™ for Spontaneously Breathing Patients

## **NICU Pocket Guide**





## **Patient Selection**



# Patient presents with one or more of the following symptoms:

- Hypoxemia
- Retractions
- Tachypnea
- Mild apnea and bradycardia
- Grunting
- Nasal flaring
- · Difficulty weaning from Nasal CPAP
- Difficulty weaning from mechanical ventilation

## Diagnoses



These symptoms are indicative of but not solely attributed to:

- Infant Respiratory
   Distress Syndrome (RDS)\*
- Bronchopulmonary Dysplasia (BPD)
- Prematurity
- Congenital Heart Defects
- Congenital diaphragmatic hernia (CDH)
- Transient Tachypnea of the Newborn (TTN)
- Meconium aspiration
- Persistent Pulmonary Hypertension (PPHN)

\*Randomized Clinical Trials show efficacy equivalent to nCPAP and NiPPV for primary support of RDS<sup>1,2</sup>

<sup>1.</sup> Kugleman et al, "A randomized pilot study comparing heated humidified high-flow nasal cannulae with NIPPV for RDS", Pediatric Pulmonology, 2014 Mar 12; 50(6) 576-83. (Clinical Trial, Prospective, Randomized, Single Site, n=76).

Lavizzari et al, "Heated, humidified high-flow nasal cannula vs nasal continuous positive airway pressure for respiratory distress syndrome of prematurity – a randomized clinical noninferiority trial", JAMA Pediatrics. 2016 Aug 8. (Clinical Trial, Prospective, Randomized, Single Site, n=316).

#### **Hi-VNI Cannula Selection**

### Fitting the Hi-VNI Cannula:

- Make sure NOT to occlude greater than 50% of the nares.
- Hi-VNI cannula prongs should be wide enough not to pinch the nasal septum (erosion risk).
- The SOLO is a single prong Hi-VNI cannula that can be used in premature, neonatal, and infant. The single prong design is as effective as a dual prong Hi-VNI cannula. The single prong simplifies NG tube placement.
- Flow rates in neonates are 1-8 L/min.

Flow Range	Tip OD
1-8 L/min	1.5 mm
1-8 L/min	1.5 mm
1-8 L/min	1.9 mm
1-8 L/min	1.9 mm
1-8 L/min	1.9 mm
1-20 L/min	1.9 mm
5-40 L/min	2.7 mm
5-40 L/min	4.8 mm
	1-8 L/min 1-8 L/min 1-8 L/min 1-8 L/min 1-8 L/min 1-8 L/min 1-20 L/min 5-40 L/min

## **Hi-VNI Cannula Application:**

- Only Vapotherm Hi-VNI cannulae should be used with the Vapotherm Precision Flow®
- · Select the appropriate Hi-VNI cannula based on the above sizing chart
- Place the Hi-VNI cannula on the patient before attaching the delivery tube
- Allow the system to reach the set point (temperature display will stop flashing) before connecting delivery tube to the Hi-VNI cannula
- The Precision Flow's operational L/min range is locked depending on the disposable patient circuit (DPC) selected:
  - PF-DPC-HIGH (Blue packaging): 5-40 L/min
  - PF-DPC-LOW (Red packaging): 1-8 L/min



## **Therapy Implementation and Maintenance**



STARTING L/MIN

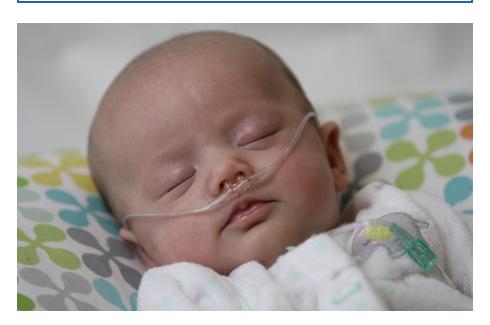
The recommended starting flow rate is 4-6 L/min, even in VLBW infants<sup>3,4</sup>. Titrate to clinical effect to maximum of 8 L/min as needed.



Set temperature to 36-37°C.



Start and titrate FiO<sub>2</sub> as needed to achieve target SpO<sub>2</sub>.



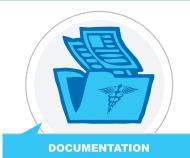
- 3. Yoder et al, "Consensus approach to nasal high-flow therapy in neonates", Journal of Perinatology. (2017) 00, 1-5.
- 4. McQueen et al, "Safety and long term outcomes with high flow nasal cannula therapy in neonatology: a large retrospective cohort study", Journal of Pulmonary Respiratory Medicine. 2014 Dec; 4(6): 216. (Clinical Trial, Retrospective, Not Randomized, Multicenter, Cohort Analysis, n=1363).

## **Monitoring Therapy**



#### **Patient Parameters:**

- Indices of work of breathing (WOB)
- SpO<sub>2</sub>
- · PCO,
- FiO<sub>2</sub>
- Nasopharynx patency
- · Feeding tolerance



#### **Documentation:**

#### **Patient**

- Heart rate
- Respiratory rate
- Work of breathing (WOB)
- · SpO<sub>2</sub>

#### Device

- Flow rate
- FiO<sub>2</sub>
- Temperature
- Water level
- Cannula size

## Weaning

# Weaning L/min

Wean in 0.5-1 L/min increments as patient tolerates

Consider further wean titrated on clinical assessment of work of breathing

If at less than 4 L/min you see rainout, consider dropping temperature to no lower than 34°C

Assess for further wean and/or discontinuation

Conventional cannula or room air



WEAN BY L/MIN OR FIO<sub>2</sub>

Vapotherm Hi-VNI **Technology parameters** (L/min & FiO<sub>2</sub>) are independent of each other. Adjustment of L/min will impact work of breathing while adjustment of FiO maintains patient SpO<sub>3</sub>. Monitoring patients' response to each change requires continuous assessment of breath sounds, respiratory rate, physical characteristics (e.g nasal flaring, grunting and retractions).

## Weaning FiO,

Return FiO<sub>2</sub> to range acceptable for SpO<sub>2</sub> requirement

Patient assessment of HR, RR, SpO<sub>2</sub>

Continue FiO<sub>2</sub> wean to maintain SpO<sub>3</sub> targets

## **Aerosol Medication and Specialty Gases**

#### **Use with Aerosol Medication**

Treating patients with respiratory disorders frequently requires combined use of Hi-VNI Technology with aerosolized medication. For practice considerations to do so, refer to the "Aerosol Delivery with HVNI Pocket Guide" and the "Aerosol Medication Delivery with HVNI Therapy Practice Summary."

#### **Use with Nitric Oxide**



- Vapotherm Hi-VNI Technology is verified for use with multiple nitric oxide delivery systems. To confirm your system is compatible with Vapotherm, contact your local representative.
- Vapotherm Nitric Oxide Disposable Patient Circuits (DPCs):

PF-NODPC-LOW 1-8 L/min PF-NODPC-HIGH 5-40 L/min

 Note: Refer to the Instructions for Use provided with your nitric oxide system and with the Nitric Oxide circuit.

#### Use with Precision Flow Heliox®

- Vapotherm offers an ideal solution for convenient delivery of conditioned helium-oxygen gas mixtures (Heliox).
- Heliox has a significantly lower density than typical air/oxygen mixtures.
- The lower gas density reduces the work of breathing by reducing the force needed to move gas through the airways.
- Heliox is commonly used on patients with diseases of increased airway resistance, such as bronchiolitis, asthma, post-extubation stridor, airway compression, intra and extrathoracic airway obstruction.
- Precision Flow Heliox strategies follow the same general clinical guidelines for air-oxygen mixtures, except FiO<sub>2</sub> should be titrated between 0.21 and 0.4 since higher oxygen concentrations (and lower helium concentrations) would result in a less significant clinical effect.
- Standard Vapotherm Disposable Patient Circuits (DPCs) may be used with the Precision Flow Heliox.

PF-DPC-LOW 1-8 L/min PF-DPC-HIGH 5-40 L/min 20 40

