# **Newborn Critical Care Center (NCCC) Clinical Guidelines**

# **Treatment of Hypotension for Extremely Low Birth Weight Infants**

#### **BACKGROUND**

In extremely low birth weight infants (ELBWs), hypotension within the first 24 hours after birth is most commonly caused by the premature infant's inability to properly transition to extra-uterine life, marked by immaturity of the myocardium. Hypotension in otherwise well ELBWs is associated with low systemic afterload and a large PDA, but not left ventricular dysfunction.[1] Other contributing factors may include anemia, adrenal insufficiency, and less commonly, hypovolemia. The incidence of hypotension increases with decreasing gestational age.

The literature on early use of vasopressors in ELBWs and its association with morbidity and mortality is predominantly observational and demonstrates mixed results. Use of inotropes has been associated with IVH, BPD, NEC, ROP, and death.[2,3] Treatment with anti-hypotensive medications within the first 24 hours of life has also been associated with an increased risk of death and neurodevelopmental impairment at 18-22 months adjusted age, even when adjusted for infant's clinical illness, [4] and an increased risk of sensorineural hearing loss.[5] In contrast, treatment of isolated hypotension has been associated with a higher rate of survival without major morbidity and a lower rate of severe cerebral abnormalities when compared to no treatment in risk-matched groups.[6] The Management of Hypotension in Preterm Infants (HIP) randomized controlled trial attempted to determine whether restricting the use of inotropes after diagnosis of low blood pressure in the first 72 hours of life affects survival without significant brain injury. This study was terminated early due to significant issues with enrollment and thus underpowered; the study did not detect differences in clinical outcomes between groups treated with dopamine vs placebo.[7]

#### **DEFINITION OF HYPOTENSION**

Uncertainty persists regarding the 'normal' values for blood pressure and which treatment thresholds should be used. As a general guideline, within the first 72 hours, the lowest value for 'normal' mean arterial pressure (MAP) will be equivalent to the gestational age (GA) in weeks (ex: for a 25 week infant ~ normal MAP 25 mmHg). However, these are general guidelines and should always be considered in the context of other potential clinical indicators of systemic hypotension.

# **EVALUATION**

Central blood pressure monitoring (i.e., an umbilical arterial line) is the gold standard and generally treatment decisions should be based on these UAC measurements rather than cuff blood pressure measurements. Assessment of cardiac contractility and filling volumes by echocardiography may also assist in decision-making.[8]

### **MEDICAL TREATMENT OPTIONS**

**Dobutamine** stimulates an increase in heart rate (chronotropy) and contractility (inotropy), allowing for more effective pumping. The increase in cardiac output makes dobutamine a good choice for neonates with myocardial dysfunction.

**Dopamine** increases the mean arterial pressure to improve peripheral and renal perfusion. Decreasing pulmonary vascular resistance makes dopamine an optimal choice when treating hypotension associated with persistent pulmonary hypertension.

#### ALGORITHM DURING THE FIRST 24 HOURS OF LIFE

- If the MAP is less than GA minus 5 mmHg (or, if 22 weeks, less than 18 mmHg) in asymptomatic infants or there is evidence of hypoperfusion (prolonged cap refill, lactic acidosis, etc.) – consider treatment with a single bolus of lactated ringer solution or normal saline (10 mL/kg). Also consider blood loss and the need for blood products for volume expansion. If the blood pressure does not increase, see below for recommended vasopressor treatment.
- 2. In asymptomatic infants, if the MAP is between GA minus 5 mmHg (or, if 22 weeks, 18 mmHg) and the normal MAP based on GA: may observe for at least one hour before considering treatment.
  - a. If the MAP is rising by at least 1 mmHg/hr, may continue to observe.
  - b. If the MAP is not increasing at this rate, consider treatment with a single bolus of lactated ringer solution or normal saline or blood products (as described above); if unresponsive, see below for recommended vasopressor treatment.

#### **Vasopressor Treatment:**

- Begin vasopressor treatment with dobutamine or dopamine at an infusion rate of 5 mcg/kg/min; increase in increments of 2 mcg/kg/min every 15 minutes (consider increasing more aggressively for severe hypotension) until a MAP > GA is achieved or a dose of 15 mcg/kg/min is reached.
- 2. If MAP < GA on a single vasopressor agent infusion of 15 mcg/kg/min, begin a second agent (dopamine or dobutamine) infusion at 5 mcg/kg/min; increase in increments of 2 mcg/kg/min every 15 minutes (consider increasing more aggressively for severe hypotension) until a MAP > GA is achieved or a dose of 15 mcg/kg/min is reached.
- 3. If MAP < GA on 15 mcg/kg/min of both dobutamine and dopamine, administer a test dose of hydrocortisone. If the MAP increases within 6 hours, begin continuing doses of hydrocortisone. Please refer to the Hydrocortisone Stress Dose Guidelines.
- 4. If the MAP does not respond to these therapies, consider echocardiography to further characterize the pathology.
- If MAP > GA, treatment is not indicated unless accompanied by objective evidence of shock. In this case, first and second line vasopressor agents may be increased to 20 mcg/kg/min each.

6. Infrequently, an epinephrine infusion may be indicated in the ELBW population if MAP remains < GA on infusions of dopamine and dobutamine (each at 20 mcg/kg/min) as well as stress dose steroids. If used, the starting dose for an epinephrine infusion would be 0.05 mg/kg/min.

# **ALGORITHM AFTER THE FIRST 24 HOURS OF LIFE**

1. If MAP < GA, begin vasopressor treatment. Fluid administration is not indicated unless there is evidence of volume depletion.

# **Vasopressor Treatment:**

- Begin vasopressor treatment with dobutamine or dopamine at an infusion rate of 5 mcg/kg/min; increase in increments of 2 mcg/kg/min every 15 minutes until a MAP > GA is achieved or a dose of 15 mcg/kg/min is reached. Consider increasing more aggressively for severe hypotension.
- 2. If MAP < GA on a single vasopressor agent infusion of 15 mcg/kg/min, begin a second agent (dopamine or dobutamine) infusion at 5 mcg/kg/min; increase in increments of 2 mcg/kg/min every 15 minutes until a MAP > GA is achieved or a dose of 20 mcg/kg/min is reached. Consider increasing more aggressively for severe hypotension.
- 3. If the MAP does not respond to these therapies, consider echocardiography to further characterize the pathology.
- 4. If MAP < GA on 20 mcg/kg/min of both dobutamine and dopamine, administer a test dose of hydrocortisone. If the MAP increases within 6 hours, begin continuing doses of hydrocortisone. Please refer to the Hydrocortisone Stress Dose Guidelines.

#### References:

- Aldana-Aguirre JC, Deshpande P, Jain A, Weisz DE. Physiology of Low Blood Pressure During the First Day After Birth Among Extremely Preterm Neonates. J Pediatr. 2021 Sep;236:40-46.e3. doi: 10.1016/j.jpeds.2021.05.026. Epub 2021 May 18. PMID: 34019882.
- 2. Wong, JJ et al. Inotrope use among extremely preterm infants in Canadian neonatal intensive care units: variation and outcomes. Am J Perinatol. 2015; 1: 32.
- Zaveri PG, Walker AM, Upadhyay K, Talati AJ. Use of Vasopressors in Extremely Preterm Infants in First Week of Life. Am J Perinatol. 2023 Apr;40(5):513-518. doi: 10.1055/s-0041-1729558. Epub 2021 May 14. PMID: 33990125.
- 4. Batton, B. et al. Early blood pressure, antihypotensive therapy and outcomes at 18-22 months' corrected age in extremely preterm infants. Arch Dis Child Fetal Neonatal Ed. 2016; 3: 101.
- Gogcu S, Washburn L, O'Shea TM. Treatment for hypotension in the first 24 postnatal hours and the risk of hearing loss among extremely low birth weight infants. J Perinatol. 2020 May;40(5):774-780. doi: 10.1038/s41372-020-0628-y. Epub 2020 Feb 26. PMID: 32103159; PMCID: PMC7185479.
- Durrmeyer X. et al. Abstention or intervention for isolated hypotension in the first 3 days of life in extremely
  preterm infants: association with short-term outcomes in the EPIPAGE 2 cohort study. Arch Dis Child Fetal
  Neonatal Ed. 2017; 102:F490-496.
- Dempsey EM, Barrington KJ, Marlow N, O'Donnell CPF, Miletin J, Naulaers G, Cheung PY, Corcoran JD, El-Khuffash AF, Boylan GB, Livingstone V, Pons G, Macko J, Van Laere D, Wiedermannova H, Straňák Z; HIP consortium. Hypotension in Preterm Infants (HIP) randomised trial. Arch Dis Child Fetal Neonatal Ed. 2021 Jul;106(4):398-403. doi: 10.1136/archdischild-2020-320241. Epub 2021 Feb 24. PMID: 33627329; PMCID: PMC8237176.
- Goldsmith JP, Keels E. Recognition and Management of Cardiovascular Insufficiency in the Very Low Birth Weight Newborn. Pediatrics. 2022 Mar 1;149(3):e2021056051. doi: 10.1542/peds.2021-056051. PMID: 35224636.

Reviewed April 2024 – Orgel MD / Patterson MD