



NEONATAL OPIOID WITHDRAWAL SYNDROME (NOWS)

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Learning Objectives

- Define neonatal opioid withdrawal syndrome
- Apply knowledge of signs and symptoms of NOWS to recognize these infants early
- Demonstrate general understanding of non-pharmacologic vs. pharmacologic management indications
- Recall the long term effects of NOWS and utilize this information to care for these infants long-term
- Educate families on clinical symptoms, management, and potential complications of NOWS

Neonatal Opioid Withdrawal Syndrome (NOWS) Key Points:

1. Incidence and relevance of NOWS
 - a. Coinciding with the Opioid epidemic starting in the early 2000s, we have seen a concomitant 5x increase in the incidence of NOWS
 - b. Approximately 7.3 out of 1000 infants are affected by NOWS
 - c. Maternal opioid use and NOWS disproportionately affect low-income women and those less likely to seek prenatal care
2. Etiology of NOWS
 - a. Opioids are natural and synthetic substances that activate mu-opioid receptors in the central nervous system and gastrointestinal tract. Substances include morphine, codeine, heroin, methadone, fentanyl, hydromorphone, and buprenorphine.
 - b. Neonatal opioid withdrawal syndrome refers to antenatal exposure of infants to opioid and the subsequent withdrawal they experience after delivery.
 - c. Most infants born with NOWS have been exposed to opioids within at least one week of delivery.
3. Clinical Signs & Symptoms of NOWS
 - a. Symptoms of NOWS affects multiple organ systems
 - b. NOWS is a result of the dysregulation of one or more of four domains
 - c. The domains can be remembered using the acronym, SOMA:
 - i. Sleep – fragmented sleep, shorter sleep cycles, difficulty maintaining an alert state between sleep cycles
 - ii. Overstimulation – irritability, tachypnea, excessive crying, difficulty feeding, poor weight gain, adverse GI issues (vomiting, diarrhea, excessive gassiness)
 - iii. Movement – jitters, tremors, hypertonicity, seizure activity
 - iv. Autonomic dysfunction – excessive yawning, sweating, fever, sneezing, mottling, nasal congestion



4. Differential diagnosis
 - a. The differential is very broad and largely dependent on the domains affected in each individual patient
 - b. Common differential diagnoses:
 - i. Hypoglycemia, hypocalcemia, other metabolic abnormalities
 - ii. Hypoxic-ischemic encephalopathy
 - iii. Transient tachypnea of newborn, respiratory distress syndrome, pulmonary hypertension, congenital heart disease, congenital pneumonia
 - iv. Infectious process, including TORCH infections and STDs
5. NOWS work-up and diagnosis
 - a. Ideally, mothers will be followed by OBGYNs and their opioid status will be known prior to delivery
 - b. Diagnosis is primarily clinical, utilizing history, maternal or infant toxicology testing, and infant signs & symptoms
 - c. Finnegan scoring system can be used to evaluate the clinical likelihood an infant is presenting with NOWS
 - d. Gold standard: meconium sample
 - i. Pros: longer window of exposure for identifying toxins
 - ii. Cons: more invasive and time-intensive; can take days for result to return
 - e. Urine toxicology
 - i. Pros: easy to obtain, quick result
 - ii. Cons: shorter window for identifying toxins
 - f. Umbilical cord testing
 - i. Pros: easy to obtain at time of delivery
 - ii. Cons: efficacy of test reliability is variable
6. Finnegan Scoring System
 - a. Lists 21 symptoms frequently observed in opioid-exposed infants
 - b. Each symptom and its degree of severity are scored, and scores are totaled as the final Finnegan score
 - c. Used to guide treatment plans and track progression/regression throughout clinical course
 - d. Provides a framework for when pharmacological management may be necessary
 - e. Additional scoring systems include the Mother Opioid Treatment: Human Experimental Research (MOTHER) and Eat, Sleep, Console (ESC) amongst others
 - f. Use of scoring system improves clinical outcomes and reduces length of hospital stay (scoring system used is institution preference)
7. Non-pharmacologic management of infants with NOWS
 - a. Individualized to infant's specific clinical signs and symptoms
 - b. Interventions are targeted at reducing stimuli that may aggravate patient symptoms:
 - i. Dim lighting
 - ii. Quiet environment
 - iii. Keeping infant with mother
 - iv. Skin-to-skin contact
 - v. Swaddling
 - vi. Breastfeeding with lactation consultation if needed
 - vii. Infant positioning
8. Pharmacologic management of infants with NOWS
 - a. First-line options: morphine, methadone, buprenorphine
 - i. Choice is largely physician and institution dependent



- ii. Combination of morphine (short-acting) and methadone (long-acting) may be beneficial to gain rapid control and maintenance of symptoms
 - b. Adjunctive therapies: phenobarbital, clonidine
 - i. Phenobarbital is used for its sedative and anticonvulsant properties and is also useful for polysubstance exposure of opioids with barbiturates or benzodiazepines.
 - ii. Clonidine can lessen symptoms of withdrawal, including tachycardia, fever, sweating, sneezing, and yawning.
 - c. Infants are monitored for a minimum of 72 hours and weaned off medication prior to discharge.
9. Potential long-term effects and longitudinal care for infants with NOWS
 - a. Infants are at an increased risk for:
 - i. Vision problems
 - ii. Motor abnormalities
 - iii. Behavioral delays
 - iv. Cognitive delays
 - v. Poor weight gain and failure to thrive
 - vi. Otitis media
 - vii. Child abuse/neglect
 - viii. Sudden unexpected infant death (SUID)
 - ix. Future substance abuse
 - b. Follow-up with pediatrician or primary care provider should be scheduled within 48 hours of hospital discharge
 - c. Additional provider follow up:
 - i. Pediatric ophthalmology

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