Persistent Wheeze in Infants: A Guide for General Pediatricians

Yolanda Yu, DO; Charles R. Esther, Jr., MD, PhD; Clement L. Ren, MD, MBA; and Ceila Loughlin, MD

ABSTRACT

Infants with persistent wheeze is a common diagnostic challenge for the general pediatrician because of the broad differential diagnoses. The initial diagnostic approach should include a comprehensive history, physical examination, and chest radiography. Additional testing may be warranted. Involvement of a pediatric pulmonary subspecialist may also be indicated. [Pediatr Ann. 2019;48(3):e110-e114.]

Wheeze is one of the most common respiratory symptoms in the pediatric population. The Tucson Children’s Respiratory Study (TCRS), a longitudinal birth cohort study of healthy full-term infants, demonstrated that 34% of children had at least one episode of wheeze before age 3 years and 50% of children wheezed in the first 6 years of life.1 Similarly, a large prospective, population-based longitudinal birth cohort study in Sweden found that 25% of children wheezed before age 2 years.2

Several risk factors have been implicated in recurrent or persistent wheeze, typically defined as three or more episodes of wheeze in 1 year (Table 1).1,3 Although multiple risk factors are associated with persistent and recurrent wheeze, the “Estudio Internacional de Sibilancias en Lactantes [International Study of Wheezing in Infants]” (EISL), a population-based study in Latin America and Europe, showed that nursery school attendance and colds during the first 3 months of life were the most important risk factors.3 The EISL also found that breast-feeding for more than 3 months was a protective factor against wheezing.3

DIAGNOSIS AND MANAGEMENT

The initial history should include a detailed account of the wheezing episodes, including onset, frequency, duration, and associated palliative and provoking factors. Birth history (eg, prematurity, respiratory distress at birth), medical history (eg, eczema, allergic rhinitis), family history (eg, atopy, asthma), and environmental history (eg, cigarette smoke exposure, day care attendance) are also important to review. Physical examination should include evaluation of respiratory rate and oxygen saturation, in addition to a thorough respiratory assessment.4

With regards to diagnostic testing, an initial chest radiograph with posteroanterior and lateral views is recommended.4 Further diagnostic testing may not be needed for infants whose wheeze responds well to commonly used therapies such as bronchodilators and inhaled corticosteroids. However, as outlined below, more advanced testing may be needed for infants who do not respond to conventional therapy.5

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DIFFERENTIAL DIAGNOSES

The most common causes of persistent and recurrent wheeze include infections, asthma, allergies, and gastroesophageal reflux disease. However, the differential for infantile wheeze is broad and includes protracted bacterial bronchitis, anatomic abnormalities, swallowing dysfunction, and others. A referral to a pediatric pulmonary subspecialist may be helpful to help establish the diagnosis and facilitate any needed testing.

INFECTIONS

Viral infections are one of the most common causes of wheeze in infancy. The TCRS demonstrated that in the first year of life, the prevalence of wheezing with lower respiratory illnesses was 32%. The TCRS also showed that this was 17.3% in the second year of life and 12% in the third year of life. Risk factors for wheezing with lower respiratory illnesses include maternal smoking and maternal age.

Asthma is one of the most common etiologies of persistent wheezing in children. It is also the most common chronic medical disease in the pediatric population and a leading cause of hospitalization. Uncontrolled asthma results in significant morbidity, mortality, and health care costs in children. The diagnosis of asthma in infants can be challenging, particularly when trying to distinguish it from viral-mediated wheeze. Because there is no test available that can easily diagnose asthma, the diagnosis is often made based on symptom history, family history, and physical examination findings. Symptoms characteristic of asthma include wheeze, nonproductive cough, shortness of breath, and activity limitation. Adjunctive clinical tests can also be helpful in diagnosis. The Global Initiative for Asthma guidelines note the following to be useful diagnostic aids: therapeutic trial of inhaled corticosteroids and as needed short-acting beta2-agonists, atopy testing, chest radiography, lung function testing (in older children), measuring exhaled nitric oxide, and asthma predictive indices.

The Asthma Predictive Index (API) was developed to aid in the early identification of children at high risk for developing asthma. It has been shown that 68% of school-aged children with a negative API did not develop asthma, whereas 76% of children with a positive API had symptoms of asthma at least once between ages 6 and 13 years. Since the API was developed, other predictive models have been created. The modified Asthma Predictive Index (mAPI) is one such clinical tool. The mAPI is for use in children younger than age 3 years and with more than four wheezing episodes per year. A study demonstrated that a positive mAPI increased the probability of future asthma diagnosis. It should be noted that all of these indices were created to predict future asthma and not the clinical response to asthma therapies in infants.

If the clinician suspects asthma and respiratory symptoms are frequent or

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**TABLE 1.**

<table>
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<th>Risk Factors for Persistent Wheeze</th>
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<td>Nursery school attendance</td>
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<td>Rhinitis apart from colds</td>
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<tr>
<td>Colds during the first 3 months of life</td>
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<tr>
<td>Eczema during the first year of life</td>
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<tr>
<td>Maternal smoking</td>
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<td>Mold stains on the walls of the household</td>
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<td>Hispanic ethnic background</td>
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<td>Maternal asthma</td>
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<td>Male gender</td>
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Adapted from Martinez et al. and Garcia-Marcos et al.
uncontrolled, current guidelines recommend initiating controller therapy. Asthma treatment in children should follow a stepwise approach. Inhaled short-acting beta2-agonists are used for symptomatic relief. When starting asthma controller therapy, an inhaled corticosteroid is the preferred initial treatment. Response to therapy should be closely monitored. After 3 months of well-controlled symptoms, a step-down in treatment may be considered. Consider a step-up in therapy, if symptoms are persistent 4 to 6 weeks after beginning treatment, but only after differential diagnosis, medication adherence, and correct inhaler technique are reviewed. If further increases in asthma therapy are considered, a referral to a pediatric asthma specialist is recommended.

ALLERGIES

Allergies are one consideration in persistent wheeze in infancy. The Generation R Study found no significant association between early introduction of cow’s milk, peanuts, soy, gluten, and hen’s egg with wheezing at any age. The American Thoracic Society (ATS) clinical practice guidelines do not recommend empiric food avoidance in “infants without eczema who have persistent wheeze despite treatment with standard therapies.”

GASTROESOPHAGEAL REFLUX DISEASE

Gastroesophageal reflux disease (GERD) is a common cause of persistent wheezing. In a study of children younger than age 2 years with recurrent and persistent wheezing, GERD contributed to wheeze in 37% of patients. Although wheeze is one common symptom of GERD, other associated symptoms include cough, feeding refusal, discomfort, dysphagia, failure to thrive, Sandifer syndrome, recurrent regurgitation, stridor, and hoarseness.

Recent clinical guidelines from ATS recommend 24-hour esophageal pH monitoring for “infants with persistent wheeze that is not relieved by bronchodilators, inhaled corticosteroids, or systemic corticosteroids.” Studies such as barium contrast studies, ultrasonography, esophago-gastro-duodenoscopy, and scintigraphy are not recommended for the diagnosis of GERD in infants. Previously empiric trials of anti-reflux therapy were given but this is no longer advised in infants. Instead, the Pediatric Gastroesophageal Reflux Clinical Practice Guidelines suggest thickening feeds and adjusting feeding frequency and volume to avoid overfeeding. If these interventions are unsuccessful, consider a 2- to 4-week trial of amino-acid–based or extensively hydrolyzed protein-based formula. If nonpharmacologic treatments fail to improve GERD, referral to a pediatric gastroenterologist is recommended.

PROTRACTED BACTERIAL BRONCHITIS

Another cause of wheezing in infancy is protracted bacterial bronchitis (PBB). This diagnosis is characterized by the presence of a chronic wet or productive cough for greater than 4 weeks that resolves with a 2- to 4-week course of an appropriate oral antibiotic. Although chronic cough is the defining symptom in PBB, wheezing has also been reported. ATS guidelines also recommend bronchoscopy with bronchoalveolar lavage (BAL) in infants with persistent wheeze despite standard therapies. Common bronchoscopic findings in children with PBB include airway malacia, bronchitis, and purulent secretions. The most common bacterial pathogens detected in BAL cultures from children diagnosed with PBB include Haemophilus influenzae, Streptococcus pneumoniae, and Moraxella catarrhalis. Management of PBB is typically a 2- to 4-week course of an appropriate oral antibiotic, although the 2008 British Thoracic Society guidelines suggested a 4- to 6-week course of antibiotics and chest physiotherapy for children with PBB.

SWALLOWING DYSFUNCTION

Swallowing dysfunction is another cause of persistent wheezing in infancy.
Sheikh et al. described 13 neurologically normal infants with chronic respiratory symptoms secondary to aspiration. Their respiratory symptoms resolved once their aspiration resolved. Studies have reported that 10% to 15% of infants with respiratory symptoms but not a chronic illness have swallowing dysfunction.

Other signs and symptoms of aspiration include coughing, choking and/or gagging during feeding, apnea, bradycardia, recurrent pneumonia or bronchitis, and failure to thrive. Recent ATS guidelines recommend swallow function study in “infants without neurologic pathology with persistent wheeze that is not relieved by bronchodilators, inhaled corticosteroids or systemic corticosteroids.” Treatment often includes changes to the feeding regimen.

FOREIGN BODY ASPIRATION

Although rare, foreign body aspiration can be one explanation for persistent wheeze. One study showed that wheezing was a common symptom and physical examination finding in children hospitalized with suspected foreign body aspiration. Chest radiography should be obtained in children with suspected foreign body aspiration. Management is rigid bronchoscopic removal of the aspirated foreign body.

ANATOMIC ABNORMALITIES

Anatomic abnormalities can cause persistent wheezing, although this is more prevalent in infants who have failed standard therapies. In a study of 603 children with chronic respiratory symptoms who underwent bronchoscopic evaluation, 34 (5.6%) had tracheomalacia and/or bronchomalacia. Other anatomic abnormalities that may present with wheeze include vascular rings, vascular slings, and airway compression by vascular structures. ATS recommends flexible bronchoscopy in infants with persistent wheeze who have failed conventional therapies. If airway evaluation reveals tracheomalacia and/or bronchomalacia, management is often observation, although occasional intervention is indicated. If a vascular ring, a vascular sling, or airway compression due to vascular structures is seen, surgical correction is often performed on symptomatic patients.

CYSTIC FIBROSIS

Cystic fibrosis (CF) is a rare cause of wheezing in infants. It is a life-limiting autosomal recessive disorder resulting from mutation of the CF transmembrane conductance regulator (CFTR) gene. Clinical presentation varies depending on the CF genotype, but can include failure to thrive, recurrent respiratory infections, and decreased lung function. The majority of new CF diagnoses are the result of positive newborn screen (NBS). Although NBS allows for early diagnosis, confirmation of the diagnosis with a sweat chloride test is still recommended. A positive sweat chloride test will demonstrate elevated sweat chloride concentration indicative of a CFTR defect. Children with CF should be treated at an accredited CF center.

PRIMARY CILIARY DYSKINESIA

Primary ciliary dyskinesia (PCD) is another rare cause of wheezing in infants. PCD is an autosomal recessive disorder characterized by motile ciliary dysfunction. Characteristic findings include daily wet productive cough, daily rhinosinusitis, history of neonatal respiratory distress syndrome, and organ laterality defects. Chronic otitis media with effusion is also often seen in PCD. Diagnostic testing includes nasal nitric oxide measurement, genetic testing, and electron microscopy of ciliary ultrastructure. Children with PCD should be managed at a PCD foundation clinical center or by a multidisciplinary team at an accredited CF center.

CONCLUSION

In summary, persistent wheezing in infants is a commonly encountered complaint with multiple etiologies. Although respiratory viral infections and asthma represent the most common etiologies, other diagnoses should be considered for infants who do not respond to standard therapies. A detailed history, thorough physical examination, and chest radiography will help narrow the differential diagnoses. Further evaluation should be tailored to the likely etiology. Consultation with a pediatric pulmonary subspecialist may also be warranted depending on severity of respiratory symptoms and suspected etiology.

REFERENCES

7. Wright AL, Holberg C, Martinez FD, Taussig LM. Relationship of parental smoking to