

# Community-Acquired Pneumonia in Infants and Children

MICHAEL OSTAPCHUK, M.D., DONNA M. ROBERTS, M.D., and RICHARD HADDY, M.D.  
*University of Louisville School of Medicine, Louisville, Kentucky*

Community-acquired pneumonia is one of the most common serious infections in children, with an annual incidence of 34 to 40 cases per 1,000 children in Europe and North America. When diagnosing community-acquired pneumonia, physicians should rely mainly on the patient's history and physical examination, supplemented by judicious use of chest radiographs and laboratory tests as needed. The child's age is important in making the diagnosis. Pneumonia in neonates younger than three weeks of age most often is caused by an infection obtained from the mother at birth. *Streptococcus pneumoniae* and viruses are the most common causes in infants three weeks to three months of age. Viruses are the most frequent cause of pneumonia in preschool-aged children; *Streptococcus pneumoniae* is the most common bacterial pathogen. *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* often are the etiologic agents in children older than five years and in adolescents. In very young children who appear toxic, hospitalization and intravenous antibiotics are needed. The symptoms in outpatients who present with community-acquired pneumonia can help determine the treatment. Knowing the age-specific causes of bacterial pneumonia will help guide antibiotic therapy. Childhood immunization has helped decrease the incidence of invasive *Haemophilus influenzae* type B infection, and the newly introduced heptavalent pneumococcal vaccine may do the same for *Streptococcus pneumoniae* infections. (Am Fam Physician 2004;70:899-908. Copyright© 2004 American Academy of Family Physicians.)

See page 801 for definitions of strength-of-recommendation labels.

The term "community-acquired pneumonia" (CAP) refers to a pneumonia in a previously healthy person who acquired the infection outside a hospital. CAP is one of the most common serious infections in children, with an incidence of 34 to 40 cases per 1,000 children in Europe and North America.<sup>1-3</sup> Although death from CAP is rare in industrialized countries, lower respiratory tract infection is one of the leading causes of childhood mortality in developing countries.<sup>4,5</sup>

## Etiology

Determining the cause of pneumonia in a child is often difficult, but the patient's age can help narrow the list of likely etiologies. Table 1<sup>6-9</sup> lists common and less common causes of CAP by age group.

Group B streptococcus and gram-negative enteric bacteria are the most common pathogens in neonates (i.e., birth to 20 days) and are obtained via vertical transmission

from the mother during birth. Anaerobic organisms may be acquired from chorioamnionitis. Pneumonia in infants aged three weeks to three months is most often bacterial; *Streptococcus pneumoniae* is the most common pathogen.

In infants older than four months and in preschool-aged children, viruses are the most frequent cause of CAP; respiratory syncytial virus (RSV) is the most common. Viral pneumonia occurs more often in the fall and winter than in the spring and summer. Bacterial infections can occur at any time of the year in preschool- and school-aged children and in adolescents.

*S. pneumoniae* is the most common bacterial cause of CAP after the neonatal period. Less common bacterial etiologies include *Haemophilus influenzae* type B, *Moraxella catarrhalis*, and *Staphylococcus aureus*. *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* frequently are associated with CAP in preschool-aged children and are common causes



**TABLE 1**  
**Causes of Community-Acquired Pneumonia by Age Group**

Age	Common causes	Less common causes
Birth to 20 days	Bacteria <i>Escherichia coli</i> Group B streptococci <i>Listeria monocytogenes</i>	Bacteria Anaerobic organisms Group D streptococci <i>Haemophilus influenzae</i> <i>Streptococcus pneumoniae</i> <i>Ureaplasma urealyticum</i> Viruses Cytomegalovirus Herpes simplex virus
3 weeks to 3 months	Bacteria <i>Chlamydia trachomatis</i> <i>S. pneumoniae</i> Viruses Adenovirus Influenza virus Parainfluenza virus 1, 2, and 3 Respiratory syncytial virus	Bacteria <i>Bordetella pertussis</i> <i>H. influenzae</i> type B and nontypeable <i>Moraxella catarrhalis</i> <i>Staphylococcus aureus</i> <i>U. urealyticum</i> Virus Cytomegalovirus
4 months to 5 years	Bacteria <i>Chlamydia pneumoniae</i> <i>Mycoplasma pneumoniae</i> <i>S. pneumoniae</i> Viruses Adenovirus Influenza virus Parainfluenza virus Rhinovirus Respiratory syncytial virus	Bacteria <i>H. influenzae</i> type B <i>M. catarrhalis</i> <i>Mycobacterium tuberculosis</i> <i>Neisseria meningitis</i> <i>S. aureus</i> Virus Varicella-zoster virus
5 years to adolescence	Bacteria <i>C. pneumoniae</i> <i>M. pneumoniae</i> <i>S. pneumoniae</i>	Bacteria <i>H. influenzae</i> Legionella species <i>M. tuberculosis</i> <i>S. aureus</i> Viruses Adenovirus Epstein-Barr virus Influenza virus Parainfluenza virus Rhinovirus Respiratory syncytial virus Varicella-zoster virus

Information from references 6-9.

of CAP in older children and adolescents.<sup>10,11</sup> Pertussis should be considered in all children with CAP, especially if immunizations are not current. *Mycobacterium tuberculosis* also may cause CAP in children at risk for exposure. Coinfection with two or more microbial agents is more common than previously thought, with a rate of up to 41 percent in hospitalized patients.<sup>6</sup>

## Clinical Evaluation

The strongest predictors of pneumonia in children are fever, cyanosis, and more than one of the following signs of respiratory distress: tachypnea, cough, nasal flaring, retractions, rales, and decreased breath sounds.<sup>5,12,13</sup>

Pneumonia should be suspected if tachypnea occurs in a patient younger than two years with a temperature



higher than 38°C (100.4°F). Measurement of tachypnea requires a full one-minute count while the child is quiet. The World Health Organization's age-specific criteria for tachypnea are the most widely used: a respiratory rate of more than 50 breaths per minute in infants two to 12 months of age; more than 40 breaths per minute in children one to five years of age; and more than 30 breaths per minute in children older than five years.<sup>14</sup>

Children without fever or symptoms of respiratory distress are unlikely to have pneumonia.<sup>1,15</sup>

Key points of the clinical assessment are given in Table 2.<sup>9</sup> The patient history, taken at the time of diagnosis, should include the child's age, immunization status,

hospitalizations, day care attendance, and recent exposures, travel, and antibiotic use.

The physician should review the child's history to identify any underlying cardiac or pulmonary diseases, immune deficiencies, or neuromuscular disorders. Inquires should be made about possible foreign object aspiration or ingestion of toxic substances. Findings not related to the respiratory tract, such as lethargy, poor feeding, vomiting, diarrhea, abdominal pain, irritability, and signs of dehydration, also should be noted.

The physical examination begins with an overall assessment of the child's well-being and identification of obvious signs of hypoxia and dehydration. The child (especially a younger child) is checked for "toxic

TABLE 2

## Clinical Assessment of Community-Acquired Pneumonia in Children

<i>Clinical clue</i>	<i>Suggested diagnosis or interpretation</i>
<b>History</b>	
Day care attendance	Viral infection
Exposure to infectious diseases	Viral or Mycoplasma infection, tuberculosis
Hospitalization	Nosocomial infection
Missing immunizations	Pneumococcal or Haemophilus influenzae infection, pertussis
Antibiotic therapy within previous month	Infection with resistant bacterial strain
Recent travel	Fungal infection, influenza, severe acute respiratory syndrome
<b>Physical examination</b>	
Elevated temperature	Pneumonia is unlikely without fever and more than one respiratory sign.
Respiratory signs	In patients with respiratory signs and no fever, consider reactive airway disease, aspiration of foreign body, chemical ingestion, or an underlying cardiac or pulmonary disorder.
Grunting	
Nasal flaring	
Rales	
Retractions	
Tachypnea	
Use of accessory muscles for breathing	
Wheezing	
<b>Laboratory studies*</b>	
Complete blood count	Not helpful in distinguishing etiology
Erythrocyte sedimentation rate	Not helpful in distinguishing etiology
C-reactive protein level	Not helpful in distinguishing etiology
Gram stain and culture	Helpful if specimen is adequate
Polymerase chain reaction	Helpful with Mycoplasma and Chlamydia infections
Rapid viral antigen testing	Useful if available
Serologies	Not helpful in acute settings
<b>Imaging</b>	
Chest radiograph*	Not helpful in distinguishing etiology

\*—Not routinely recommended.

Information from reference 9.



appearance,” tachypnea, elevated temperature, retractions, grunting, and use of accessory muscles. The upper respiratory tract should be examined for evidence of otitis media, rhinorrhea, nasal polyposis, and pharyngitis. Physical signs such as heart murmurs or nail clubbing may suggest underlying cardiac or pulmonary disease.

Older children and adolescents are more likely to have findings such as rales, dullness to percussion, bronchial breath sounds, tactile fremitus, and a pleural rub.<sup>7</sup> Careful auscultation with an appropriate-sized stethoscope may reveal localized rales and wheezing in younger children. Children with dehydration may have no abnormal auscultatory findings.

## Diagnostic Testing

In most children with CAP, identification of the causative organism is not critical.<sup>16</sup> Patients with severe symptoms, those who are hospitalized, and those who have a complicated clinical course should undergo diagnostic testing to determine the etiology. The cause also should be determined if there appears to be a community outbreak.

Older children and adolescents may be able to produce sputum for Gram stain and culture. Adequate specimens contain more than 25 leukocytes and fewer than 25 squamous epithelial cells per low-power field.<sup>7</sup>

Rapid antigen tests are available for RSV, parainfluenza 1, 2, and 3, influenza A and B, and adenovirus. These assays, which are performed on specimens collected from the nasopharynx, can help determine the etiology of viral pneumonia.<sup>1,15,17</sup> Nasopharyngeal specimens for bacterial culture or antigen assays are less useful, because bacteria commonly colonize on the nasopharynx.<sup>1,15,17</sup> Antigen and antibody assays for pneumococcal infection are not sensitive enough to be helpful in diagnosing *S. pneumoniae* infection. In the future, detection of pneumococcal immune complexes may offer a rapid etiologic diagnosis in children older than two years.<sup>18</sup>

Serologic testing for IgM or an increase in IgG titers may be performed for *Mycoplasma* and *Chlamydia* species. However, serologic tests often provide only a retrospective diagnosis and are more useful in establishing the causative agent during an outbreak than in treating individual children.<sup>19</sup> Cultures for *Mycoplasma* and *Chlamydia* are not routinely recommended. Polymerase chain reaction testing is not readily accessible, and positive results do not necessarily imply causality.<sup>16</sup>

The complete blood count with differential, C-reactive protein level, and erythrocyte sedimentation rate do not differentiate bacterial from viral infection and

should not be measured routinely.<sup>1,20</sup> After the neonatal period, the incidence of bacteremia in children is so low that collection of blood for cultures can be individualized to patients who appear more ill.<sup>6</sup>

Chest radiographs do not differentiate causative agents of CAP as well as was previously thought.<sup>1,21</sup> Lobar consolidation classically has been associated with pneumococcal infections, and interstitial infiltrates have been associated with viral infections. However, both lobar consolidation and interstitial infiltrates have been identified in all types of infections—viral alone, bacterial alone, and viral-bacterial.<sup>22</sup> Chest radiographs should not be obtained routinely in children with mild, uncomplicated lower respiratory tract infection.<sup>1,13</sup> Indications for chest radiographs include ambiguous clinical findings, prolonged pneumonia, pneumonia that is unresponsive to antibiotic therapy, and the possibility of complications such as pleural effusions.<sup>13</sup>

Oxygen saturation should be assessed by pulse oximetry in children with respiratory distress, significant tachypnea, or pallor.<sup>15</sup>

## Treatment

Treatment decisions are based on the child's age and clinical and epidemiologic factors.<sup>1,8</sup> Antibiotic therapy should be initiated promptly in children who are thought to have bacterial CAP. Because definitive information about the causative organism is usually unknown, the choice of antibiotic is empiric.<sup>16</sup> Table 3 lists the recommended outpatient and inpatient antibiotic therapies for different age groups, based on evidence-based guidelines from the University of Cincinnati Children's Hospital, the Alberta Medical Association, and the British Thoracic Society.<sup>1,8,13,15,23</sup>

### INPATIENTS

Table 4 lists factors to consider in deciding whether inpatient management is necessary.<sup>1,15</sup> Hospitalization is required for all infants from birth to 20 days of age, infants three weeks to three months of age with fever, and all children who appear toxic. Hospital admission criteria for children four months to five years of age include hypoxemia or a respiratory rate of more than 70 breaths per minute. Other indicators for admission include difficulty breathing, intermittent apnea, grunting, poor feeding, and inadequate observation or supervision by family.<sup>1</sup> Admission criteria for older children include hypoxemia, cyanosis, a respiratory rate of more than 50 breaths per minute, difficulty breathing, and inadequate observation or supervision by family.<sup>1</sup>



## NEONATES

Infants younger than three weeks with respiratory distress always should be admitted to a hospital, and a diagnosis of bacterial pneumonia should be assumed until proved otherwise. Cultures of blood, urine, and cerebrospinal fluid should be obtained, and treatment with ampicillin and gentamicin, with or without cefotaxime (Claforan), should be started as soon as possible.<sup>7,8,23</sup>

## INFANTS

Infants three weeks to three months of age who are suspected of having bacterial pneumonia require immediate attention, particularly if they are febrile, tachypneic, or appear toxic.<sup>7</sup> These patients are best treated in a hospital. Initial therapy consists of cefuroxime (Ceftin) or cefotaxime.<sup>15</sup> Blood, urine, and cerebrospinal fluid cultures; a complete blood count with differential; and a chest radiograph should be obtained.<sup>7,8</sup> Once stabilized, infants may be changed to an oral antibiotic for 10 days.

*Chlamydia trachomatis* infection should be suspected in infants who are afebrile or nontoxic and have a dry cough.<sup>7,8</sup> These patients often have a peripheral eosinophilic pleocytosis.<sup>7</sup> In such cases, treatment guidelines recommend outpatient treatment with an oral macrolide and close follow-up.<sup>8</sup> If the physician decides to treat the infant as an inpatient, intravenous erythromycin is the drug of choice.<sup>8</sup>

## PRESCHOOL-AGED CHILDREN

Viruses cause most cases of pneumonia in preschool-aged children (i.e., four months to five years of age).<sup>1,9,16,24</sup> Although most physicians start antibiotic therapy, guidelines allow for withholding treatment if a viral etiology is suspected and close follow-up can be ensured.<sup>1,9,13,15,16,24</sup> These children usually have associated symptoms of viral infection, such as pharyngitis, rhinorrhea, and diarrhea (Table 5).<sup>1</sup>

Pneumococcal infection is the most common cause of bacterial pneumonia in this age group.<sup>7,24</sup> The onset of pneumococcal pneumonia is usually abrupt, with none of the prodromal symptoms associated with viral illnesses. A child suspected of having pneumococcal pneumonia who is not hypoxic, in distress, or unstable can be treated empirically with high-dosage amoxicillin.<sup>8,9,13,24</sup> The physician may initiate therapy with a single dose of ceftriaxone (Rocephin).<sup>13</sup> Alternatives include amoxicillin-clavulanic acid (Augmentin), azithromycin (Zithromax), cefaclor (Ceclor), clarithromycin (Biaxin), and erythromycin.

Preschool-aged children who require hospital admission are treated with cefuroxime or cefotaxime.<sup>7,15</sup> Once

the child is afebrile and stable, he or she is switched to an oral antibiotic and treated on an outpatient basis.

## OLDER CHILDREN

*S. pneumoniae* is a significant pathogen in school-aged children and adolescents (i.e., five to 18 years of age) with CAP. *M. pneumoniae* and *C. pneumoniae* infections also are more common in these children than in other age groups.<sup>1</sup>

In school-aged children, pneumococcal pneumonia usually begins with a high fever and sputum-producing cough. *M. pneumoniae* infection often begins with headache or gastrointestinal symptoms; rhinorrhea is uncommon.<sup>7</sup> Other symptoms, such as fever, arthralgia, and cough, in a school-aged child suggest Mycoplasma infection.<sup>1</sup> A clinical decision tree for ruling out Mycoplasma infection in children with CAP is presented in Figure 1.<sup>25</sup>

CAP caused by *C. pneumoniae* routinely begins with pharyngitis, followed by a cough and high fever. If pneumococcal pneumonia is suspected, high-dosage amoxicillin may be used.<sup>1,23,26</sup> If *M. pneumoniae* or *C. pneumoniae* infection is suspected, a macrolide antibiotic is the drug of choice.<sup>1,23,26</sup> Azithromycin, erythromycin, or clarithromycin may be used as a single agent in this age group because all of these agents provide adequate coverage for penicillin-sensitive pneumococcus.<sup>1,7,13</sup> Therapeutic options for hospitalized patients are cefuroxime or cefotaxime in addition to a macrolide.<sup>15,26</sup>

### Determining Risk of *Mycoplasma pneumoniae* Infection

*The rightsholder did not grant rights to reproduce this item in electronic media. For the missing item, see the original print version of this publication.*

Figure 1.



TABLE 3

**Therapeutic Management of Community-Acquired Pneumonia**

<i>Patient age</i>	<i>Outpatient</i>	<i>Inpatient</i>	<i>Critically ill</i>
Birth to 20 days	Admit	<p>Ampicillin IV or IM:  Age &lt;7 days:  Weight &lt;2 kg (4.4 lb): 50 to 100 mg per kg per day in divided doses every 12 hours  Weight ≥2 kg: 75 to 150 mg per kg per day in divided doses every 8 hours  Age ≥7 days:  Weight &lt;1.2 kg (2.6 lb): 50 to 100 mg per kg per day divided every 12 hours  Weight 1.2 to 2 kg: 75 to 150 mg per kg per day in divided doses every 8 hours  Weight &gt;2 kg: 100 to 200 mg per kg per day in divided doses every 6 hours  <i>plus</i>  Gentamicin IV or IM:  ≥37 weeks of gestation  <i>and</i>  Age zero to 7 days: 2.5 mg per kg every 12 hours  Age &gt;7 days: 2.5 mg per kg every 8 hours  <i>with or without</i>  Cefotaxime (Claforan) IV:  Age ≤7 days:  100 mg per kg per day in divided doses every 12 hours  Age &gt;7 days:  150 mg per kg per day in divided doses every 8 hours</p>	<p>Ampicillin IV or IM, in same dosages as for inpatients  <i>plus</i>  Gentamicin IV or IM, with or without cefotaxime IV, in same dosages as for inpatients</p>
3 weeks to 3 months	<p>If patient is <b>afebrile</b>:  Azithromycin (Zithromax), 10 mg per kg orally on day 1, then 5 mg per kg per day on days 2 through 5  <i>or</i>  <b>Erythromycin</b>, 30 to 40 mg per kg per day orally in divided doses every 6 hours for 10 days  Admit if patient is febrile or hypoxic.</p>	<p><b>Erythromycin</b>, 40 mg per kg per day IV in divided doses every 6 hours*  <b>If patient is febrile, add</b> one of these agents:  <b>Cefotaxime</b>, 200 mg per kg per day IV in divided doses every 8 hours*  <i>or</i>  <b>Cefuroxime</b> (Ceftin), 150 mg per kg per day IV in divided doses every 8 hours*</p>	<p><b>Cefotaxime</b>, 200 mg per kg per day IV in divided doses every 8 hours  <i>plus</i> <b>cloxacillin</b> (Tegopen), 150 to 200 mg per kg per day IV in divided doses every 6 hours*  <i>or</i>  <b>Cefuroxime</b> alone, 150 mg per kg per day IV in divided doses every 8 hours*</p>

*continued*



TABLE 3 (continued)

Patient age	Outpatient	Inpatient	Critically ill
4 months to 5 years	<p>Amoxicillin, 90 mg per kg per day orally in divided doses every 8 hours for 7 to 10 days</p> <p>Consider initial dose of ceftriaxone (Rocephin), 50 mg per kg per day IM, up to 1 g per day. Follow with oral therapy for full course.</p> <p>Alternatives: amoxicillin-clavulanic acid (Augmentin), azithromycin, cefaclor (Ceclor), clarithromycin (Biaxin), erythromycin</p>	<p>Cefotaxime, 150 mg per kg per day IV in divided doses every 6 hours*</p> <p>or</p> <p>Cefuroxime, 150 mg per kg per day IV in divided doses every 8 hours*</p> <p>If the patient has pneumococcal infection: Ampicillin alone, 200 mg per kg per day IV in divided doses every 8 hours*</p>	<p>Cefuroxime, 150 mg per kg per day IV in divided doses every 8 hours, plus erythromycin, 40 mg per kg per day IV or orally in divided doses every 6 hours for 10 to 14 days*</p> <p>or</p> <p>Cefotaxime, 200 mg per kg per day IV in divided doses every 8 hours, plus cloxacillin, 150 to 200 mg per kg per day IV in divided doses every 6 hours for 10 to 14 days</p>
5 years and older	<p>Azithromycin, 10 mg per kg (maximum of 500 mg) orally on day 1, followed by 5 mg per kg per day on days 2 through 5</p> <p>or</p> <p>Clarithromycin, 15 mg per kg per day orally in divided doses every 12 hours for 7 to 10 days</p> <p>or</p> <p>Erythromycin, 40 mg per kg per day orally in divided doses every 6 hours for 7 to 10 days</p> <p>If the patient has pneumococcal infection: Amoxicillin alone, 90 mg per kg per day orally in divided doses every 8 hours</p>	<p>Cefuroxime, 150 mg per kg per day IV in divided doses every 8 hours</p> <p>plus</p> <p>Erythromycin, 40 mg per kg per day IV or orally in divided doses every 6 hours for 10 to 14 days</p> <p>If pneumococcal infection is confirmed: Ampicillin alone, 200 mg per kg per day IV in divided doses every 8 hours</p>	<p>Cefuroxime, 150 mg per kg per day IV in divided doses every 8 hours</p> <p>plus</p> <p>Erythromycin, 40 mg per kg per day IV or orally in divided doses every 6 hours for 10 to 14 days</p>

IV = intravenous; IM = intramuscular.

\*—Duration is total length of IV and oral therapy. Consider switching to oral therapy when child has no complications and is afebrile, clinically improving, not experiencing diarrhea, and tolerating oral intake.

Information from references 1, 8, 13, 15, and 23.

#### DURATION OF THERAPY

Antibiotic therapy should be continued for seven to 10 days in patients with uncomplicated CAP, although no controlled study of the optimal treatment duration exists.<sup>15</sup> Follow-up for outpatients should be done at 24 to 72 hours

after diagnosis.<sup>1,3,24</sup> Reevaluation is necessary in children who continue to have unresolved symptoms or fever at 48 hours after diagnosis. In these patients, physicians should suspect inappropriate antibiotic therapy or a lung complication, such as an empyema or abscess.<sup>1</sup>



Asymptomatic children with normal physical findings after treatment do not need follow-up chest radiographs<sup>7,24,27</sup>

Repeat chest radiographs or computed tomographic scans are recommended if the illness is protracted or a complication such as empyema is suspected.<sup>1</sup>

Penicillin-resistant pneumococcus is a concern in the treatment of CAP. Evidence indicates that inpatient intravenous therapy with a penicillin or cephalosporin is effective against penicillin-resistant pneumococcus. Oral  $\beta$ -lactam antibiotics are appropriate first-line therapies for patients with ambulatory CAP.<sup>8,16,28,29</sup>

### Prevention

Childhood immunizations have helped greatly in the prevention of pneumonia in children. Pneumonia is a known complication of rubeola, varicella, and pertussis. These illnesses and the pneumonias related to them rarely are seen today because of routine childhood immunizations.<sup>30</sup> Pneumonia caused by *H. influenzae* type B also is rare, because of routine administration of the Hib vaccine.

In February 2000, a new heptavalent pneumococcal vaccine was licensed for use in the United States. This

### The Authors

MICHAEL OSTAPCHUK, M.D., is assistant professor of family and geriatric medicine and associate director of the family practice residency program at the University of Louisville (Ky.) School of Medicine. Dr. Ostapchuk received his medical degree from the University of Kentucky College of Medicine, Lexington, and a master's degree in science education from the University of Southern California, Los Angeles. He completed a pediatric residency at East Carolina University School of Medicine, Greenville, N.C., and a family medicine residency at the University of Louisville School of Medicine.

DONNA M. ROBERTS, M.D., is director of predoctoral education and associate professor in the Department of Family and Geriatric Medicine at the University of Louisville School of Medicine. Dr. Roberts received her medical degree from West Virginia University School of Medicine, Morgantown, and completed a family medicine residency at the University of Louisville School of Medicine.

RICHARD HADDY, M.D., is professor and vice chairman for academic affairs in the Department of Family and Geriatric Medicine at the University of Louisville School of Medicine. He received his medical degree from Michigan State University College of Human Medicine, East Lansing, and completed a family medicine residency at the University of Iowa Hospitals and Clinics, Iowa City. Dr. Haddy also completed a one-year fellowship in infectious diseases at the Saginaw campus of Michigan State University College of Human Medicine.

Address correspondence to Michael Ostapchuk, M.D., Department of Family and Geriatric Medicine, University of Louisville School of Medicine, Med Center One, 501 East Broadway, Suite 240, Louisville, KY 40202 (e-mail: m0osta01@gwise.louisville.edu). Reprints are not available from the authors.

TABLE 4

#### Considerations for Admission of Patients with Community-Acquired Pneumonia

*The rightsholder did not grant rights to reproduce this item in electronic media. For the missing item, see the original print version of this publication.*

vaccine produces immunity for the seven most common disease-producing serotypes of *S. pneumoniae* in children. Widespread use of this vaccine is expected to decrease the incidence of invasive pneumococcal disease dramatically.<sup>8,31</sup> Initial trials have shown the vaccine to be highly effective in preventing invasive disease.<sup>31</sup> There appears to be a similar decrease in the incidence of pneumococcal pneumonia alone,<sup>32</sup> although the study

TABLE 5

#### Characteristics of Viral and Bacterial Pneumonia

*The rightsholder did not grant rights to reproduce this item in electronic media. For the missing item, see the original print version of this publication.*



## Strength of Recommendations

Key clinical recommendations	Label	Reference
Childhood immunizations, including the recently released heptavalent pneumococcal vaccine, help decrease the incidence of invasive pneumococcal disease.	A	32
The complete blood count, C-reactive protein level, and erythrocyte sedimentation rate do not help determine the etiology of CAP. Organism-specific testing may be used in some instances.	B	20
The chest radiograph is not helpful in differentiating the causative agents of CAP in children.	B	21
The patient's age and associated symptoms can guide the physician in determining the etiology of CAP in children.	C	1
Empiric treatment for CAP in children is based on the patient's age and symptoms.	C	1

CAP = community-acquired pneumonia.

that demonstrated this finding was limited by the inability to definitively diagnose the cause of pneumonia. More recent longitudinal data have shown a significant decrease in rates of invasive pneumococcal disease, particularly in children two years of age and younger.<sup>33,34</sup>

Vaccination with the heptavalent pneumococcal vaccine may increase carriage of noninvasive serotypes. A recent study<sup>35</sup> suggests that immunized children are more likely to develop otitis media with serotypes not covered by the heptavalent vaccination. However, the long-term benefits of heptavalent pneumococcal vaccine are promising and continue to be investigated.

The American Academy of Pediatrics (AAP) recommends influenza vaccination for all high-risk children six months of age and older.<sup>36</sup> To protect against the complications of influenza, including pneumonia, the AAP also recommends vaccination of all children six months through 23 months of age, to the extent logistically and economically feasible.<sup>36,37</sup>

Members of various family medicine departments develop articles for "Practical Therapeutics." This article is one in a series coordinated by the Department of Family and Geriatric Medicine at the University of Louisville School of Medicine, Louisville, Ky. Guest editor of the series is James G. O'Brien, M.D.

The authors indicate that they do not have any conflicts of interest. Sources of funding: none reported.

The authors thank Maggie Steptoe for assistance in the preparation of the manuscript.

## REFERENCES

- British Thoracic Society Standards of Care Committee. British Thoracic Society guidelines for the management of community acquired pneumonia in childhood. *Thorax* 2002;57(suppl 1):i1-24.
- Murphy TF, Henderson FW, Clyde WA Jr, Collier AM, Denny FW. Pneumonia: an eleven-year study in a pediatric practice. *Am J Epidemiol* 1981;113:12-21.
- Jokinen C, Heiskanen L, Juvonen H, Kallinen S, Karkola K, Korppi M, et al. Incidence of community-acquired pneumonia in the population of four municipalities in eastern Finland. *Am J Epidemiol* 1993;137:977-88.
- Boschi-Pinto C, Debay M. Informal consultation on epidemiologic estimates for child health. 11-12 June 2001. Accessed online February 27, 2004, at: [http://www.who.int/child-adolescent-health/New\\_Publications/Overview/Report\\_of\\_CHERG\\_meeting.htm](http://www.who.int/child-adolescent-health/New_Publications/Overview/Report_of_CHERG_meeting.htm).
- Redd SC, Vreuls R, Metsing M, Mohobane PH, Patrick E, Moteetee M. Clinical signs of pneumonia in children attending a hospital outpatient department in Lesotho. *Bull World Health Organ* 1994;72:113-8.
- Juven T, Mertsola J, Waris M, Leinonen M, Meurman O, Roivainen M, et al. Etiology of community-acquired pneumonia in 254 hospitalized children. *Pediatr Infect Dis J* 2000;19:293-8.
- Gaston B. Pneumonia. *Pediatr Rev* 2002;23:132-40.
- McIntosh K. Community-acquired pneumonia in children. *N Engl J Med* 2002;346:429-37.
- Bradley JS. Management of community-acquired pediatric pneumonia in an era of increasing antibiotic resistance and conjugate vaccines. *Pediatr Infect Dis J* 2002;21:592-8, 613-4.
- Principi N, Esposito S. *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* cause lower respiratory tract disease in paediatric patients. *Curr Opin Infect Dis* 2002;15:295-300.
- Heiskanen-Kosma T, Korppi M, Jokinen C, Kurki S, Heiskanen L, Juvonen H, et al. Etiology of childhood pneumonia: serologic results of a prospective, population-based study. *Pediatr Infect Dis J* 1998;17:986-91.
- Margolis P, Gadomski A. The rational clinical examination. Does this infant have pneumonia? *JAMA* 1998;279:308-13.
- Cincinnati Children's Hospital Medical Center Health Policy and Clinical Effectiveness Program. Evidence based clinical practice guideline. Community acquired pneumonia in children 60 days to 17 years of age. Accessed online February 27, 2004, at: <http://www.cincinnatichildrens.org/NR/rdonlyres/E25BBF59-5EDD-4B75-A28F-24B6F6C35016/0/PneumoniaGL.pdf>.
- World Health Organization. Essential drugs and medicines policy. Drugs used in bacterial infections. Accessed online February 27, 2004, at: [http://www.who.int/medicines/library/bacterial\\_model\\_pres/bacterial\\_content.shtml](http://www.who.int/medicines/library/bacterial_model_pres/bacterial_content.shtml).
- Alberta Clinical Practice Guidelines Steering Committee. Guideline for the diagnosis and management of community acquired pneumonia: pediatric. 2002. Accessed online June 1, 2004, at: [http://www.albertadoctors.org/bcm/ama/ama-website.nsf/AllDocSearch/87256DB000705C3F87256E0500553605/\\$File/PNEUMONIA\\_PEDIATRICS.PDF](http://www.albertadoctors.org/bcm/ama/ama-website.nsf/AllDocSearch/87256DB000705C3F87256E0500553605/$File/PNEUMONIA_PEDIATRICS.PDF).



## Community-Acquired Pneumonia

16. McCracken GH Jr. Diagnosis and management of pneumonia in children. *Pediatr Infect Dis J* 2000;19:924-8.
17. Korppi M, Heiskanen-Kosma T, Leinonen M, Halonen P. Antigen and antibody assays in the aetiological diagnosis of respiratory infection in children. *Acta Paediatr* 1993;82:137-41.
18. Korppi M, Leinonen M. Pneumococcal immune complexes in the diagnosis of lower respiratory infections in children. *Pediatr Infect Dis J* 1998;17:992-5.
19. Principi N, Esposito S. Emerging role of *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* in paediatric respiratory-tract infections. *Lancet Infect Dis* 2001;1:334-44.
20. Nohynek H, Valkeila E, Leinonen M, Eskola J. Erythrocyte sedimentation rate, white blood cell count and serum C-reactive protein in assessing etiologic diagnosis of acute lower respiratory infections in children. *Pediatr Infect Dis J* 1995;14:484-90.
21. Esposito S, Bosis S, Cavagna R, Faelli N, Begliatti E, Marchisio P, et al. Characteristics of *Streptococcus pneumoniae* and atypical bacterial infections in children 2-5 years of age with community-acquired pneumonia. *Clin Infect Dis* 2002;35:1345-52.
22. Korppi M, Kiekara O, Heiskanen-Kosma T, Soimakallio S. Comparison of radiological findings and microbial aetiology of childhood pneumonia. *Acta Paediatr* 1993;82:360-3.
23. Bradley JS, Nelson JD. 2002-2003 Nelson's pocket book of pediatric antimicrobial therapy. 15th ed. Philadelphia: Lippincott, Williams & Wilkins;2002:31-5.
24. Peter G. Pneumonia. In: Burg FD, Gellis SS, Kagan BM. Gellis & Kagan's current pediatric therapy. 16th ed. Philadelphia: Saunders;1999:32-5.
25. Fischer JE, Steiner F, Zucol F, Berger C, Martignon L, Bossart W, et al. Use of simple heuristics to target macrolide prescription in children with community-acquired pneumonia. *Arch Pediatr Adolesc Med* 2002;156:1005-8.
26. Nelson JD. Community-acquired pneumonia in children: guidelines for treatment. *Pediatr Infect Dis J* 2000;19:251-3.
27. Gibson NA, Hollman AS, Paton JY. Value of radiological follow up of childhood pneumonia. *BMJ* 1993;307:1117.
28. Heffelfinger JD, Dowell SF, Jorgensen JH, Klugman KP, Mabry LR, Musher DM, et al. Management of community-acquired pneumonia in the era of pneumococcal resistance: a report from the Drug-Resistant *Streptococcus pneumoniae* Therapeutic Working Group. *Arch Intern Med* 2000;160:1399-408.
29. McCracken GH Jr. Etiology and treatment of pneumonia. *Pediatr Infect Dis J* 2000;19:373-7.
30. Mandell GL, Douglas RG, Bennett JE, Dolin R. Mandell, Douglas, and Bennett's principles of practice of infectious diseases. 5th ed. Philadelphia: Churchill Livingstone, 2000:2416-7.
31. Black S, Shinefield H, Fireman B, Lewis E, Ray P, Hansen JR, et al. Efficacy, safety and immunogenicity of heptavalent pneumococcal conjugate vaccine in children. Northern California Kaiser Permanente Vaccine Study Center Group. *Pediatr Infect Dis J* 2000;19:187-95.
32. Black SB, Shinefield HR, Ling S, Hansen J, Fireman B, Spring D, et al. Effectiveness of heptavalent pneumococcal conjugate vaccine in children younger than five years of age for prevention of pneumonia. *Pediatr Infect Dis J* 2002;21:810-5.
33. Lin PL, Michaels MG, Janosky J, Ortenzo M, Wald ER, Mason EO Jr. Incidence of invasive pneumococcal disease in children 3 to 36 months of age at a tertiary care pediatric center 2 years after licensure of the pneumococcal conjugate vaccine. *Pediatrics* 2003;111(4 pt 1):896-9.
34. Whitney CG, Farley MM, Hadler J, Harrison LH, Bennett NM, Lynfield R, et al. Decline in invasive pneumococcal disease after the introduction of protein-polysaccharide conjugate vaccine. *N Engl J Med* 2003;348:1737-46.
35. Eskola J, Kilpi T, Palmu A, Jokinen J, Haapakoski J, Herva E, et al. Efficacy of a pneumococcal conjugate vaccine against acute otitis media. *N Engl J Med* 2001;344:403-9.
36. Committee on Infectious Diseases. American Academy of Pediatrics. Reduction of the influenza burden in children. *Pediatrics* 2002;110:1246-52.
37. Bridges CB, Fukuda K, Uyeki TM, Cox NJ, Singleton JA; Centers for Disease Control and Prevention, Advisory Committee on Immunization Practices. Prevention and control of influenza. Recommendations of the Advisory Committee on Immunization Practices (ACIP). *MMWR Recomm Rep* 2002;51:1-31.