



Complete Summary

GUIDELINE TITLE

Evidence based clinical practice guideline for medical management of acute otitis media in children 2 months to 13 years of age.

BIBLIOGRAPHIC SOURCE(S)

Cincinnati Children's Hospital Medical Center. Evidence based clinical practice guideline for medical management of acute otitis media in children 2 months to 13 years of age. Cincinnati (OH): Cincinnati Children's Hospital Medical Center; 2004 Oct. 16 p. [113 references]

GUIDELINE STATUS

This is the current release of the guideline.

This guideline updates a previous version: Cincinnati Children's Hospital Medical Center. Evidence-based clinical practice guideline for medical management of otitis media in children 2 months to 6 years of age. Cincinnati (OH): Children's Hospital Medical Center (CHMC); 1999. 8 p.

The guideline was reviewed for currency in August 2006 using updated literature searches and was determined to be current.

**** REGULATORY ALERT ****

FDA WARNING/REGULATORY ALERT

Note from the National Guideline Clearinghouse: This guideline references a drug(s) for which important revised regulatory information has been released.

- [September 11, 2007, Rocephin \(ceftriaxone sodium\)](#): Roche informed healthcare professionals about revisions made to the prescribing information for Rocephin to clarify the potential risk associated with concomitant use of Rocephin with calcium or calcium-containing solutions or products.

COMPLETE SUMMARY CONTENT

**** REGULATORY ALERT ****

SCOPE

METHODOLOGY - including Rating Scheme and Cost Analysis

RECOMMENDATIONS

EVIDENCE SUPPORTING THE RECOMMENDATIONS

BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS

CONTRAINDICATIONS

QUALIFYING STATEMENTS
IMPLEMENTATION OF THE GUIDELINE
INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT
CATEGORIES
IDENTIFYING INFORMATION AND AVAILABILITY
DISCLAIMER

SCOPE

DISEASE/CONDITION(S)

Acute otitis media (AOM)

GUIDELINE CATEGORY

Counseling
Diagnosis
Evaluation
Management
Treatment

CLINICAL SPECIALTY

Emergency Medicine
Family Practice
Otolaryngology
Pediatrics

INTENDED USERS

Advanced Practice Nurses
Allied Health Personnel
Health Care Providers
Nurses
Patients
Physician Assistants
Physicians

GUIDELINE OBJECTIVE(S)

- To improve the use of appropriate diagnostic criteria
- To improve the use of appropriate antibiotic therapy
- To improve symptom relief
- To avoid medical complications
- To improve parental involvement in decision-making around the management of acute otitis media

TARGET POPULATION

Children age 2 months up to 13 years of age who present with signs and symptoms of acute otitis media (AOM)

Note: Children with comorbid conditions increasing the risk or severity of otitis media, including immunodeficiencies, craniofacial or neurologic abnormalities, or sensory deficits are excluded. Children with pressure equalization (PE) tubes in place are also excluded.

INTERVENTIONS AND PRACTICES CONSIDERED

Diagnosis

1. History and physical examination
2. Pneumatic otoscopy and tympanometry
3. Acoustic reflectometry

Management/Treatment

1. Appropriate analgesia including oral agents (acetaminophen or ibuprofen) or topical ear drops (anesthetic or Naturopathic Herbal Extract Ear Drops)
2. Antibiotic therapy
 - Amoxicillin (first-line treatment)
 - Amoxicillin/clavulanate [Augmentin®]
 - Cephalosporins, including cefdinir (Omnicef®), cefprozil [Cefzil], cefuroxime (Ceftin®), and ceftriaxone (Rocephin®)
 - Macrolides/azilides, including azithromycin (Zithromax®) and clarithromycin (Biaxin®)
3. Observation option with or without safety net antibiotic prescription
4. Steroids, antihistamines, decongestants, and complementary or alternative therapies (considered, but not recommended)
5. Follow-up evaluation

Consults and Referrals

1. Referral for an audiologic evaluation
2. Referral for an otolaryngological evaluation

Education

1. Educating family on the natural history of acute otitis media and middle ear effusion, signs and symptoms of clinical deterioration, and appropriate follow-up
2. Educating family on preventable and nonpreventable risk factors

MAJOR OUTCOMES CONSIDERED

- Duration of signs and symptoms of acute otitis media (fever, discomfort)
- Risk of disease recurrence
- Hearing or speech compromise or subsequent need for insertion of tympanostomy tubes

METHODOLOGY

METHODS USED TO COLLECT/SELECT EVIDENCE

Searches of Electronic Databases

DESCRIPTION OF METHODS USED TO COLLECT/SELECT THE EVIDENCE

To select evidence for critical appraisal by the group, the citations in the American Academy of Pediatrics (AAP) Clinical Practice Guideline for Acute Otitis Media were reviewed. Additionally, the Medline, EmBase, and the Cochrane databases were searched for dates of January, 2003 through June, 2004 to generate an unrefined, combined evidence database using a search strategy focused on answering clinical questions relevant to otitis media and employing a combination of Boolean searching on human-indexed thesaurus terms (Medical Subject Headings (MeSH) headings using an OVID Medline interface) and natural language searching on words in the title, abstract, and indexing terms. The citations were reduced by: eliminating duplicates, review articles, non-English articles, and adult articles. The resulting abstracts were reviewed by a methodologist to eliminate low quality and irrelevant citations. During the course of the guideline development, additional clinical questions were generated and subjected to the search process.

August 2006 Review

A search using the above criteria was conducted for dates of July 2004 through August 2006. Sixty-five relevant articles were selected as potential future citations for the guideline. However, none of these references was determined to require changes to the 2004 version of the recommendations.

NUMBER OF SOURCE DOCUMENTS

385

METHODS USED TO ASSESS THE QUALITY AND STRENGTH OF THE EVIDENCE

Subjective Review

RATING SCHEME FOR THE STRENGTH OF THE EVIDENCE

Not applicable

METHODS USED TO ANALYZE THE EVIDENCE

Review
Review of Published Meta-Analyses
Systematic Review

DESCRIPTION OF THE METHODS USED TO ANALYZE THE EVIDENCE

Not stated

METHODS USED TO FORMULATE THE RECOMMENDATIONS

Expert Consensus

DESCRIPTION OF METHODS USED TO FORMULATE THE RECOMMENDATIONS

The recommendations contained in this guideline were formulated by an interdisciplinary working group which performed systematic and critical literature reviews, using the grading scale below under "Type of Evidence Supporting the Recommendations," and examined current local clinical practices.

During formulation of these guidelines, the team members have remained cognizant of controversies and disagreements over the management of these patients. They have tried to resolve controversial issues by consensus where possible and, when not possible, to offer optional approaches to care in the form of information that includes best supporting evidence of efficacy for alternative choices.

RATING SCHEME FOR THE STRENGTH OF THE RECOMMENDATIONS

Not applicable

COST ANALYSIS

A formal cost analysis was not performed and published cost analyses were not reviewed.

METHOD OF GUIDELINE VALIDATION

Comparison with Guidelines from Other Groups
Internal Peer Review

DESCRIPTION OF METHOD OF GUIDELINE VALIDATION

The guidelines have been reviewed and approved by clinical experts not involved in the development process, senior management, Risk Management & Corporate Compliance, the Institutional Review Board, other appropriate hospital committees, and other individuals as appropriate to their intended purposes.

RECOMMENDATIONS

MAJOR RECOMMENDATIONS

Each recommendation is followed by evidence grades (A-X) identifying the type of supporting evidence. Definitions for the types of evidence are presented at the end of the "Major Recommendations" field.

Assessment and Diagnosis

General

Signs and symptoms of acute otitis media (AOM) are often nonspecific and overlap with those of upper respiratory infections. Clinical diagnosis is especially less reliable in the child under 2 years of age. This contributes to difficulty in accurately diagnosing AOM and in evaluating results of clinical trials (Dagan et al., 2002 [S]; Froom et al., 1990 [O]; Wald, 2003 [E]).

Note: In a chart review, only 38% of pediatricians diagnoses of AOM met the Center for Disease Control (CDC) criteria for the diagnosis (Garbutt, Jeffe, & Shackelford, 2003 [O]). Inter-rater agreement for AOM diagnosis has been measured at 64% (Blomgren & Pitkaranta, 2003 [C]).

History and Physical Examination

1. It is recommended that the components to assess for AOM in the history and physical include history of acute onset of symptoms, presence of middle ear effusion (MEE), and signs and symptoms of middle ear inflammation (American Academy of Pediatrics [AAP] Subcommittee, 2004 [S]). See table below entitled "Requirements for Diagnosis of AOM."

Note 1: Accurate diagnosis of AOM, critical to the management decision, is a competency that may be improved with specific training (Rosenfeld, 2002 [O]; Steinbach & Sectish, 2002 [O]; Pichichero & Poole, 2001 [O]). See CD-ROM available in Pratt Library (Wald & Hoberman, 2002 [E]) and an interactive case module on-line with the AAP (<http://www.aap.org/otitismedia/www/>).

Note 2: A bulging, cloudy, immobile and distinctly red tympanic membrane (TM) is most helpful in the diagnosis of AOM (Rothman, Owens, & Simel, 2003 [M]; Leibovitz et al., "Can acute otitis media," 2003 [C]; Karma et al., 1989 [D]). See table below entitled "Likelihood Ratios (LR) for Clinical Signs."

Note 3: A parental report of AOM symptoms is somewhat reliable (sensitivity 71%, specificity 80%, positive predictive value 51%, likelihood ratio [LR] 3.55) (Kontiokari et al., 1998 [C]).

Note 4: Nonspecific symptoms include cough, rhinitis, poor appetite and vomiting and have likelihood ratios (LR) near 1.0 (Heikkinen & Ruuskanen, 1995 [C], Niemela et al., 1994 [C]).

Table: Requirements for Diagnosis of AOM

- | |
|---|
| <ol style="list-style-type: none">1. History of acute onset of signs and symptoms |
|---|

<p>2. Presence of MEE indicated by one of the following:</p> <ul style="list-style-type: none"> • Bulging tympanic membrane (ear drum) • Decreased mobility of tympanic membrane • Discharge from the ear (otorrhea)
<p>3. Signs and symptoms of middle ear inflammation indicated by either:</p> <ul style="list-style-type: none"> • Red tympanic membrane or • Discomfort affecting normal activity and/or sleep (earache, otalgia)

(AAP Subcommittee, 2004 [S])

Table: Likelihood Ratios (LR) for Clinical Signs*

Sign of tympanic membrane	Positive LR (95% Confidence Interval [CI])**
Bulging	51 (36-73)
Cloudy	34 (28-42)
Distinctly impaired mobility	31 (26-37)
Distinctly red	8.4 (6.7-11)
Normal color	0.2 (0.19-0.21)
Normal mobility	0.2 (0.19-0.21)

(Rothman, Owens, & Simel, 2003 [M]; Karma et al., 1989 [D])

*Likelihood ratios quantify the change in probability of AOM when a given sign or symptom is present in a specific clinical case and depend upon a starting estimate of probability. For more information, see Appendix 2 of the original guideline document for definition and use of likelihood ratios.

**95% Confidence Interval expresses the uncertainty (precision) of a measured value; it is the range of values within which we can be 95% sure that the true value lies. A study with a larger sample will generate more precise measurements, resulting in a narrower confidence interval.

2. It is recommended that pneumatic otoscopy and/or tympanometry be used to enhance accuracy when diagnosing AOM (Spiro et al., 2004 [A]; Karma et al., 1989 [D]; Brookhouser, 1998 [S]; Pelton, 1998 [S]; Jones & Kaleida, 2003 [O]; Pichichero, 2002 [O]; Pichichero & Poole, 2001 [O]).

Note 1: Pneumatic otoscopy and tympanometry measure the degree of mobility of the tympanic membrane as an indication of the presence of MEE (Jerger, 1970 [C]; Jones & Kaleida, 2003 [O]; Pichichero, 2002 [O]; Pichichero & Poole, 2001 [O]). A large, randomized controlled trial demonstrated that the appropriate use of tympanometry would reduce otitis media diagnoses by 14 to 40% (Spiro et al., 2004 [A]).

Note 2: Acoustic reflectometry is not often used nor readily available in the Cincinnati area, though the procedure is acceptable for determining the presence of MEE (Block et al., 1999 [C]; Barnett et al., 1998 [C]; Block et al., 1998 [C]; Kimball, 1998 [S]).

3. It is recommended that ear pain (otalgia) be assessed, as determined by discomfort affecting normal activity and/or sleep (Kontiokari et al., 1998 [C]; "The assessment and management," 2001 [S]).
4. It is recommended that, for patients with recurrent AOM, additional attention be paid to parental concerns about hearing loss, speech delay, or language delay (Roberts, Rosenfeld, & Zeisel, 2004 [M]).

Management

General

AOM is a disease with a high spontaneous resolution rate (78 to 80% resolve within 7 to 14 days), and routine antibiotic therapy of all children with suspected AOM results in the treatment of many children in whom there may be either modest benefit and/or modest adverse outcomes from antibiotic therapy (Glasziou et al., 2003 [M]; Marcy et al., 2001 [M]; Rosenfeld et al., 1994 [M]; Dowell et al., "Otitis media," 1998 [S]). Moreover, the decision to use antibiotics and the specific choice of antibiotics must take into account the increasing emergence of bacterial resistance (Doern et al, 1998 [C]; Jacobs et al, 2003 [O]; Mason et al., 2003 [O]).

Note: AOM may have potentially serious complications including mastoiditis, meningitis, and intracranial abscess formation. There is an increased incidence of mastoiditis in some countries which limit use of antibiotics for AOM, but a causal relationship is not fully supported by these data. Prevalence of mastoiditis in the U.S. is 0.4%, and 2,500 cases of AOM would need to be treated with antibiotics to prevent one case of mastoiditis (number needed to treat [NNT] = 2,500) (Van Zuijlen et al., 2001 [O]).

Treatment

1. It is recommended that all children with AOM who have a positive assessment for pain be treated with an appropriate analgesic (AAP Subcommittee, 2004 [S]; "The assessment and management," 2001 [S]).

Note 1: Ear pain in AOM is self-limiting and time is the greatest factor in pain reduction (Sarrell, Cohen, & Kahan, 2003 [A]). Therefore, the immediate availability of a safe and effective analgesic is more important than which

agent is used. These include oral agents (acetaminophen or ibuprofen) or topical ear drops (anesthetic or Naturopathic Herbal Extract Ear Drops) (Perrott et al., 2004 [M]; Sarrell, Cohen, & Kahan, 2003 [A]; Sarrell, Mandelberg, & Cohen, 2001 [A]; Bertin et al., 1996 [A]; Hoberman et al., "Efficacy of Auralgan," 1997 [B]; "The assessment and management," 2001 [S]).

Note 2: In patients with a perforated eardrum and/or discharge from the ear, avoid topical analgesic ear drops as their use will likely result in severe dizziness and vomiting.

2. It is recommended that treatment with a 10-day course of antibiotics be given to children less than 2 years of age with AOM (Cohen et al., 2000 [A], 1998 [A]; AAP Subcommittee, 2004 [S]).

Amoxicillin, in the dose range of 80 to 90 mg/kg/day is effective in the treatment of a first episode of AOM or for a recurrence more than 1 month since recovery from a prior episode of AOM (Rosenfeld et al., 1994 [M]; Piglansky et al., 2003 [C]; AAP Subcommittee, 2004 [S]). See table below entitled "First-Line Antibiotic Medication and Doses" for doses for this first-line therapy. In cases when the clinician has a high suspicion for concurrent conjunctivitis-otitis media syndrome, commonly caused by a beta-lactamase producing organism, it is reasonable to consider a second-line antibiotic (Wald, 1997 [S]).

For children with allergies to penicillin, or other reasons to consider alternative antibiotics, consider a second-line antibiotic. In a child less than 1 year of age with a history of a penicillin allergy, a careful review of the reported reaction is prudent. See extended list of antibiotic options, doses, and preparations in Appendix 3 (Rosenfeld et al., 1994 [M]).

Note: While it is suggested in general that children under two years of age with AOM be treated with antibiotics, it is recognized that in certain situations observation without antibiotics or with a safety-net antibiotic prescription (SNAP) may be reasonable. See table below entitled "Safety-Net Antibiotic Prescription (SNAP) Definition and Management" for SNAP definition. Local data suggest that the relapse/recurrent rate (defined in the study as a new case of AOM occurring between 7 and 60 days from initial episode) is about 3 times higher in children less than 2 years of age (34% compared to 10%) (Siegel et al., 2004 [C]). If observation or SNAP is utilized, both the clinician and the parents are advised to be aware of the higher relapse/recurrence rate in this age group, and close follow-up must be assured (Siegel et al., 2003 [C]).

Table: First-Line Antibiotic Medication and Doses

Antibiotic	Dose and frequency	Max Daily Dose
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Antibiotic	Dose and frequency	Max Daily Dose
amoxicillin	80-90 mg/kg per day, divided <ul style="list-style-type: none"> 40-45 mg/kg twice a day (BID) or 25-30 mg/kg three times a day (TID) 	2 grams/day

Table: Safety-Net Antibiotic Prescription (SNAP) Definition and Management

<ul style="list-style-type: none"> SNAP is a prescription for an appropriate antibiotic, as determined by the practitioner, <u>written to be filled only within 5 days of the office visit.</u>
<ul style="list-style-type: none"> Instruct the parent not to fill SNAP unless symptoms worsen at any time or unless symptoms do not improve during a waiting period of 48-72 hours.
<ul style="list-style-type: none"> Instruct the parent that a well-appearing child diagnosed with AOM may quickly progress to a more severe case, and to call and/or follow-up with practitioner if this occurs.

(Siegel et al., 2003 [C])

- It is recommended that in children over age 2 years with AOM and who are well-appearing, that the treatment options be discussed with the family and that the family be involved in the decision making. The options include:
 - Treatment with a SNAP to be filled after 48 to 72 hours if symptoms do not resolve with observation. See table above for SNAP definition and management.
 - Treatment with a 5-day course of antibiotics (see treatment recommendation 2 and table above entitled "First-Line Antibiotic Medication and Doses" for discussion of antibiotic selection and doses)

Note 1: There is inadequate evidence to say that antibiotic therapy is or is not beneficial to most children with AOM (Wald, 2003 [E]).

Note 2: No antibiotic prescription, with follow-up within 48 to 72 hours, is also an option in the specific case when a practitioner would like to control the observation option more closely.

Note 3: An [observation option information sheet](#) for parents is available for use through the Cincinnati Children's Hospital Medical Center (CCHMC) Health Topics Web site (New York Regional Otitis Project, 2002 [X]; Rosenfeld, 2001 [X]).

Note 4: Parental involvement in the decision to use antibiotic therapy and the use of SNAP with pain control are effective in reducing the use of antibiotics (Pshetizky, Naimer, & Shvartzman, 2003 [B]; Siegel et al., 2003 [C]).

Note 5: In a Cincinnati office-based study, the relapse/recurrence rate for all ages (new case of AOM occurring between 7 to 60 days from initial episode) was 24% in those that filled the SNAP, compared to 11% in those that did not fill the SNAP, $p < 0.025$ (Siegel et al., 2004 [C]).

4. It is recommended that children over age 2 years with AOM and with severe illness (see table below entitled "Factors to Consider for AOM Treatment") be treated with a 5-day course of antibiotics (see treatment recommendation 2 and table above entitled "First Line Antibiotic Medication and Doses") (Kozyskyj et al., 2000 [M], 1998 [M]).
5. It is recommended, for a child with a recurrence of AOM in less than 1 month from completion of antibiotic therapy from a prior episode of AOM, or for a child who has recently been on antibiotics for other reasons, that antibiotic choices other than amoxicillin be considered (Leibovitz et al., "Recurrent acute otitis media," 2003 [C]; Carlin et al., 1987 [C]; Dowell et al., "Principles of judicious use," 1998 [S]; Klein, 1998 [S]). For extended list of antibiotic options, doses, and preparations, see Appendix 3 of the original guideline document entitled "Expanded Table of Antibiotic Options, Doses and Preparations."

Note: There is no strong evidence to support prolonged or prophylactic antibiotic therapy in recurring AOM (Williams et al., 1993 [M]; Koivunen et al., 2004 [A]). Persistent MEE is common, and parents may be counseled to expect fluid to take several weeks to months to clear (Rosenfeld & Kay, 2003 [M]; AAP Subcommittee, 2004 [S]).

6. It is **not** recommended that other therapies be used in the treatment of AOM (AAP Subcommittee, 2004 [S]).

Note 1: Steroids, antihistamines, decongestants, and complementary or alternative treatments have not been documented to be efficacious in the treatment of AOM (Butler & Van Der Voort, 2002 [M]; Flynn, Griffin, & Tudiver, 2002 [M]; Barnett et al., 2000 [C]; AAP Subcommittee, 2004 [S]). Antihistamines may prolong the duration of MEE.

Note 2: Median duration of MEE was 73 days for patients on antihistamine compared to 25 days for patients on placebo ($p = 0.04$) in a randomized controlled study (Chonmaitree et al., 2003 [A]).

Note 3: It is recognized that use of complementary and alternative medicine (CAM) is common and its use is often not reported to the primary care physician (PCP) (Eisenberg et al., 1998 [O]; Spiegelblatt, 1994 [O]). The primary care physician may take the AOM visit as an opportunity to begin a respectful discussion regarding the safety and efficacy of complementary and alternative medicine with families who report its use.

Table: Factors to Consider for AOM Treatment

<ul style="list-style-type: none">• Temperature >38.6 degrees C (101.5 degrees F) in the past 48 hours
<ul style="list-style-type: none">• Symptoms suggestive of AOM for >48 hours
<ul style="list-style-type: none">• Toxic appearance
<ul style="list-style-type: none">• Tympanic membrane of the infected ear not intact
<ul style="list-style-type: none">• Another episode of AOM within past 3 months
<ul style="list-style-type: none">• Signs of impending perforation in the infected ear as judged by examining clinician
<ul style="list-style-type: none">• A coexisting bacterial infection
<ul style="list-style-type: none">• Concerns of clinician that the family would be unable to seek medical care if the child's clinical status were to worsen
<ul style="list-style-type: none">• Concerns of caregiver or clinician that the caregiver could not gain an acceptable understanding of the therapeutic plan

Follow-up: to Treatment and Observation Options

7. It is recommended, for the first or a sporadic episode of AOM, that when the initial management approach fails, the clinician reevaluate the antibiotic decision.

If symptoms worsen at any time or if symptoms do not improve during a waiting period of 48 to 72 hours of initial presentation with AOM, and reexamination continues to suggest that AOM is the appropriate diagnosis, then start amoxicillin if not already initiated or change to an alternative antibiotic if the child is already on a first line drug.

Note: Options for alternative antibiotics include:

- Amoxicillin/clavulanate: efficacy has been shown for AOM and may be used when resistance is likely (Hoberman et al., Equivalent efficacy, 1997 [A]; Dagan et al., 2001 [C])
- Ceftriaxone intramuscularly (IM): 3 consecutive daily doses is efficacious in nonresponsive AOM for children with vomiting or otherwise unable to tolerate oral dosing (Leibovitz et al., 2000 [A]).

For children with allergies to penicillin, or other reasons to consider another antibiotic choice, see extended list of antibiotic options, doses, and preparations in Appendix 3 of the original guideline document.

8. It is recommended that the clinician reevaluate the patient in 4 to 8 weeks after diagnosis, depending on existing risk factors, to document resolution or persistence of effusion (AAP Subcommittee, 2004 [S]; Local Expert Consensus [E]).

Consults and Referrals

1. It is recommended that a practitioner have a low threshold for referral for an audiologic evaluation by a pediatric audiologist if concerns around hearing, speech, or language are raised by parents, clinician, or other caregivers because of recurrent AOM (Mandel et al., 1991 [A]; Hsu, Levine, & Giebink, 1998 [C]; Teele et al., 1990 [C]; Bachmann & Arvedson 1998 [S]).
2. It is recommended that a child be referred for an otolaryngological evaluation for:
 - Recurrent AOM (history of 6 episodes over a 12-month period taking into account the severity of episodes, clustering of episodes, and persistence of otitis media with effusion)
 - Persistent otorrhea
 - Concerns about mastoiditis, or other complications of AOM
 - Perceived need for tympanocentesis and/or myringotomy (e.g., acute episode not responsive to medical therapy)
 - Abnormal audiologic evaluation

(Froom et al., 1993 [C])

Education

1. It is recommended that the family be educated regarding the natural history of AOM, signs and symptoms of clinical deterioration, and appropriate follow-up.
2. It is recommended that the practitioner discuss with the parent that persistent MEE is common, and parents may expect fluid to take several weeks to months to clear (Rosenfeld & Kay, 2003 [M]; AAP Subcommittee, 2004 [S]).
3. It is recommended that the family be educated about preventable risk factors. These include:
 - Exposure to others (especially family members) with upper respiratory tract infections (Uhari, Mantysaari, & Niemela, 1996 [M])
 - Parental smoking or other sources of second-hand smoke (Uhari, Mantysaari, & Niemela, 1996 [M]; Ilicali et al., 1999 [C])

- Daycare attendance (Uhari, Mantysaari, & Niemela, 1996 [M]; Bradley, 2003 [C])

Note: Though daycare attendance may not be preventable, options to reduce risk of AOM include delaying daycare, selecting a setting with fewer children, and/or verifying the daycare facility's hand washing practices and availability of sinks.

- Excessive pacifier use, limiting use to when the child is falling asleep (Uhari, Mantysaari, & Niemela, 1996 [M]; Niemela et al., 2000 [A])
- Breastfeeding duration less than 3 months (Uhari, Mantysaari, & Niemela, 1996 [M])
- Bottlefeeding with the child on his/her back: assure that infants are offered bottle feedings while sitting in upright positions (Tully, Bar-Haim, & Bradley, 1995 [B])

Parents may also benefit by understanding nonpreventable risk factors or common misconceptions.

- Anatomy of the eustachian tube in young children
- It is not always known why a child gets AOM.
- Allergies do not cause AOM.

Definitions:

Evidence Based Grading Scale:

A: Randomized controlled trial: large sample
 B: Randomized controlled trial: small sample
 C: Prospective trial or large case series
 D: Retrospective analysis
 E: Expert opinion or consensus
 F: Basic laboratory research
 S: Review article
 M: Meta-analysis
 Q: Decision analysis
 L: Legal requirement
 O: Other evidence
 X: No evidence

CLINICAL ALGORITHM(S)

An algorithm is provided in the original guideline document for evaluation and management of acute otitis media in children 2 months to 13 years of age.

EVIDENCE SUPPORTING THE RECOMMENDATIONS

REFERENCES SUPPORTING THE RECOMMENDATIONS

[References open in a new window](#)

TYPE OF EVIDENCE SUPPORTING THE RECOMMENDATIONS

The type of supporting evidence is classified for the recommendations (see "Major Recommendations").

Evidence Based Grading Scale:

A: Randomized controlled trial: large sample
B: Randomized controlled trial: small sample
C: Prospective trial or large case series
D: Retrospective analysis
E: Expert opinion or consensus
F: Basic laboratory research
S: Review article
M: Meta-analysis
Q: Decision analysis
L: Legal requirement
O: Other evidence
X: No evidence

BENEFITS/HARMS OF IMPLEMENTING THE GUIDELINE RECOMMENDATIONS

POTENTIAL BENEFITS

- Effective medical management of acute otitis media in children 2 months to 13 years of age
- Improved use of appropriate diagnostic criteria
- Improved use of appropriate antibiotic therapy
- Improved symptom relief
- Avoidance of medical complications
- Improved parental involvement in decision-making around the management of acute otitis media

POTENTIAL HARMS

Not stated

CONTRAINDICATIONS

CONTRAINDICATIONS

In patients with a perforated eardrum and/or discharge from the ear, avoid topical analgesic ear drops as their use will likely result in severe dizziness and vomiting.

QUALIFYING STATEMENTS

QUALIFYING STATEMENTS

These recommendations result from review of literature and practices current at the time of their formulations. This protocol does not preclude using care modalities proven efficacious in studies published subsequent to the current revision of this document. This document is not intended to impose standards of care preventing selective variances from the guidelines to meet the specific and unique requirements of individual patients. Adherence to this pathway is voluntary. The physician in light of the individual circumstances presented by the patient must make the ultimate judgment regarding the priority of any specific procedure.

IMPLEMENTATION OF THE GUIDELINE

DESCRIPTION OF IMPLEMENTATION STRATEGY

Appropriate companion documents have been developed to assist in the effective dissemination and implementation of the guideline.

IMPLEMENTATION TOOLS

Clinical Algorithm
Patient Resources
Quick Reference Guides/Physician Guides

For information about [availability](#), see the "Availability of Companion Documents" and "Patient Resources" fields below.

INSTITUTE OF MEDICINE (IOM) NATIONAL HEALTHCARE QUALITY REPORT CATEGORIES

IOM CARE NEED

Getting Better

IOM DOMAIN

Effectiveness
Patient-centeredness

IDENTIFYING INFORMATION AND AVAILABILITY

BIBLIOGRAPHIC SOURCE(S)

Cincinnati Children's Hospital Medical Center. Evidence based clinical practice guideline for medical management of acute otitis media in children 2 months to 13 years of age. Cincinnati (OH): Cincinnati Children's Hospital Medical Center; 2004 Oct. 16 p. [113 references]

ADAPTATION

Not applicable: The guideline was not adapted from another source.

DATE RELEASED

1999 (revised 2004 Oct 29; reviewed 2006 Aug)

GUIDELINE DEVELOPER(S)

Cincinnati Children's Hospital Medical Center - Hospital/Medical Center

SOURCE(S) OF FUNDING

Cincinnati Children's Hospital Medical Center

GUIDELINE COMMITTEE

Acute Otitis Media Team Members 2004

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Community Physicians: Stephen Pleatman, MD, Chair; Robert Siegel, MD

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All Team Members and Clinical Effectiveness support staff listed above have signed a conflict of interest declaration.

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FINANCIAL DISCLOSURES/CONFLICTS OF INTEREST

All Team Members and Clinical Effectiveness support staff have signed a conflict of interest declaration.

GUIDELINE STATUS

This is the current release of the guideline.

This guideline updates a previous version: Cincinnati Children's Hospital Medical Center. Evidence-based clinical practice guideline for medical management of otitis media in children 2 months to 6 years of age. Cincinnati (OH): Children's Hospital Medical Center (CHMC); 1999. 8 p.

The guideline was reviewed for currency in August 2006 using updated literature searches and was determined to be current.

GUIDELINE AVAILABILITY

Electronic copies: Available from the [Cincinnati Children's Hospital Medical Center Web Site](#).

For information regarding the full-text guideline, print copies, or evidence-based practice support services contact the Children's Hospital Medical Center Health Policy and Clinical Effectiveness Department at HPCEInfo@chmcc.org.

AVAILABILITY OF COMPANION DOCUMENTS

The following is available:

- Guideline highlights. Acute otitis media age 2 months to 13 years. Cincinnati Children's Hospital Medical Center, 2004.

Electronic copies: Available in Portable Document Format (PDF) from the [Cincinnati Children's Hospital Medical Center Web site](#).

PATIENT RESOURCES

The following are available:

- Appendix 1: resources for patient/family education purpose. Cincinnati Children's Hospital Medical Center, 2004. Electronic copies available from the [Cincinnati Children's Hospital Medical Center Web site](#).
- Ear infections and acute otitis media. Cincinnati Children's Hospital Medical Center, 2004. Electronic copies available from the [Cincinnati Children's Hospital Medical Center Web site](#).
- Types of hearing tests. Cincinnati Children's Hospital Medical Center, 2004. Electronic copies available from the [Cincinnati Children's Hospital Medical Center Web site](#).

- Managing ear infections (acute otitis media). Cincinnati Children's Hospital Medical Center, 2004. Electronic copies available from the [Cincinnati Children's Hospital Medical Center Web site](#).

Please note: This patient information is intended to provide health professionals with information to share with their patients to help them better understand their health and their diagnosed disorders. By providing access to this patient information, it is not the intention of NGC to provide specific medical advice for particular patients. Rather we urge patients and their representatives to review this material and then to consult with a licensed health professional for evaluation of treatment options suitable for them as well as for diagnosis and answers to their personal medical questions. This patient information has been derived and prepared from a guideline for health care professionals included on NGC by the authors or publishers of that original guideline. The patient information is not reviewed by NGC to establish whether or not it accurately reflects the original guideline's content.

NGC STATUS

This summary was completed by ECRI on September 20, 1999. The information was verified by the guideline developer as of November 15, 1999. This NGC summary was updated by ECRI on December 8, 2004. The information was verified by the guideline developer on January 12, 2005. This summary was updated on May 3, 2005 following the withdrawal of Bextra (valdecoxib) from the market and the release of heightened warnings for Celebrex (celecoxib) and other nonselective nonsteroidal anti-inflammatory drugs (NSAIDs). This summary was updated by ECRI on June 16, 2005, following the U.S. Food and Drug Administration advisory on COX-2 selective and non-selective non-steroidal anti-inflammatory drugs (NSAIDs). This summary was updated by ECRI Institute on October 3, 2007 following the U.S. Food and Drug Administration (FDA) advisory on Rocephin (ceftriaxone sodium).

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