**Otitis externa:** Providing relief while avoiding complications

Treatment options may be equally effective, but risks of ototoxicity vary

**Practice recommendations**

- Topical antimicrobial otic drops, supplemented by oral analgesics, provide prompt and effective therapy (B).

- Fluoroquinolones and neomycin-polymyxin B combinations have shown equal efficacy (A), but the former are preferred when trying to guard against ototoxicity or hypersensitivity (C).

- For severely painful otitis externa, consider inserting an ear wick and giving oral narcotics (C).

- Localized cellulitis requires systemic antimicrobial therapy (C).

- No compelling evidence exists on the effectiveness of astringent topical treatments for otitis externa. Also, topical astringents are often painful, which may impede adherence (B).

**Symptoms and complications to watch for**

Pain, which may be incapacitating, is the predominant complaint and the only symptom directly related to disease severity. Patients may also experience discharge, itchiness, and, in some cases, temporary hearing loss. A fluffy white exudate may signal a rare fungal infection, perhaps complicating an underlying bacterial infection.

**Bone involvement.** A rare, extremely severe form of otitis externa, known as malignant (or necrotizing) otitis externa, is caused by a *Pseudomonas* infection that invades the posterior cranial bone. This life-threatening form of otitis externa affects primarily elderly patients with diabetes who live in hot, humid environments. Malignant otitis externa usually requires hospitalization and parenteral antibiotic therapy.
Pathogens of otitis externa

Acute inflammation of the external auditory canal most often is caused by bacterial infection. Infrequently, a fungal infection may be the cause.

*Pseudomonas aeruginosa* and *Staphylococcus aureus* are the predominant bacterial pathogens associated with otitis externa. The pathogenesis involves modification of the natural antimicrobial defense mechanisms of the external ear canal. Under healthy conditions, cerumen (ear wax), which is secreted by sebaceous and apocrine glands in the external ear canal, is acidic and contains lysozyme, an antimicrobial substance. The ear canal becomes vulnerable to bacterial infection when the cerumen barrier is disrupted by scratching or scraping, or macerated by prolonged exposure to water. Disruption of the external ear’s epithelium, caused by seborrhea, eczema, or trauma, can also increase the risk of otitis externa. Because frequent swimming is the most common predisposing factor for the disease, otitis externa is commonly referred to as “swimmer’s ear.”

As may be expected, the number of cases of otitis externa increases markedly during the summer months in temperate climates. The infection is uncommon in children aged less than 2 years.

Rarely, otitis externa appears to be caused by fungal infection. Such cases are not usually differentiated on clinical grounds, except when there is appearance of a fluffy white exudate. In most cases of fungal otitis externa, the fungus appears to be a superinfection after the bacterial infection. Thus, persistent otitis externa is typically treated with 2 or 3 courses of topical antibiotics before the clinician begins to investigate more specifically for fungal superinfection.

Secondary cellulitis. Summer often ushers in several cases of another severe form of otitis externa in children, involving a secondary cellulitis of the pre- and postauricular skin structures. If an underlying mastoiditis is uncertain, consider ordering a computerized tomography scan and referring for an otolaryngologic exam. This cellulitis infection also requires aggressive broad-spectrum oral or parenteral therapy.

Elements of successful treatment

Most treated cases of otitis externa resolve completely within 5 days. In a study conducted in the Netherlands, 35 out of 98 adult patients discontinued daily activities for a median of 4 days, and the median duration of bed rest was 3 days.

However, secondary skin and soft tissue infections can slow resolution. They may also develop if treatment is delayed or ineffective. For uncomplicated otitis externa, treatment with an appropriate topical antibiotic, plus potent analgesics to relieve pain, is the preferred therapeutic approach. Choosing the optimal topical treatment, however, requires knowledge of evolving resistance among pathogens, possible hypersensitivity to neomycin, adherence factors, and cost issues.

Topical antibiotics usually sufficient. Most cases of otitis externa without significant complications are effectively treated with just topical antibiotics; though, for more severe cases, I prefer to insert an ear wick and to use oral narcotics.

In the 1970s to 1990s, topical combination preparations containing hydrocortisone and the antibiotics neomycin sulfate and polymyxin B were the mainstay of therapy for otitis externa. Approximately 5 years ago, topical fluoroquinolones became available for otitis externa. Given this range of antibiotic choices, physicians need to know the various treatment options and other modalities available.

Oral antibiotics. Additional oral or parenteral antibiotics are usually necessary for severe cases of otitis externa with secondary cellulitis or lymphadenitis. Oral antibiotics may also be necessary with concomitant disease at other sites (sinus, middle ear) or complications.

Pain relief. Nonsteroidal anti-inflammatory drugs (NSAIDs—eg, ibuprofen), acetaminophen, and narcotics (eg, codeine) are adjunctive measures for reducing ear pain.

Enabling compliance. Patient compliance, according to one systematic review, is inversely proportional to the prescribed number of doses per day. Adherence with twice-daily dosing was significantly better than with 4-times-daily dosing for oral and topical medications. Twice-daily dosing is accepted as readily as once-daily dosing.
Most treated cases of otitis externa resolve completely within 5 days

**Is there a place for astringents?**

Before antibiotic availability, the mainstay of treatment was topical compounds containing astringents such as aluminum acetate solution (Burrow’s Solution USP, Domeboro, and others) and acetic acid solutions (Vosol Otic). These preparations are still widely used but principally to prevent otitis externa.

Topical 8% aluminum acetate solution was as effective as a commonly used antimicrobial-corticosteroid topical mixture (polymyxin-neomycin-hydrocortisone) for 25 adult patients with otitis externa (72% vs 76%, respectively). In a double-blind, randomized study involving 65 adults and children, the effectiveness of aluminum acetate equaled that of gentamicin otic solution.

In a Dutch study evaluating otitis externa in 213 adults, topical antibiotics were more effective than topical acetic acid alone and alleviated symptoms faster than acetic acid plus steroids.

The lack of a large, well-structured, placebo-controlled study or other compelling evidence casts doubt on the relative effectiveness of astringent topical treatments for otitis externa. Also, topical astringents are often painful, which may impede adherence.

Practitioners usually reserve nonspecific therapy such as acetic acid and Burrow’s solution for prevention and self-medication for mild symptoms. Studies that include measures of clinical goals, such as pain relief, cessation of symptoms, and eradication of infection, are needed before astringent agents can be recommended for routine therapy of moderate to severe otitis externa. Most treatment data concerning otitis externa have centered on therapy with antimicrobial otic drops.

**Preparing the ear for antibiotic administration**

Topical antibiotics are presumed to eradicate pathogens most effectively through direct contact at concentrations exceeding the minimum inhibitory concentration (MIC$_{90}$).

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**FIGURE 1**

**Ear wick within the external ear canal**

The ear wick is inserted into the external auditory canal until the remaining edge is flush with the external ear. In smaller children, one fourth to one third of the wick may be cut to prevent it from falling out prematurely. Several drops of topical antibiotics are instilled onto the wick until it has fully expanded. At home, a few drops of topical antibiotic should be applied to the wick every few hours for the first 24 hours to keep the wick moist. The wick should be removed with tweezers within 24 to 48 hours if it has not already fallen out.
Fluffy white exudate may signal fungal infection, perhaps complicating a bacterial infection.

Once the edema subsides and much of the debris in the external ear canal has been cleared, topical antibiotics can be instilled.

**Debride if possible.** Though the severity of pain may preclude debridement, removing debris from the external ear canal during the office visit will maximize delivery of the medication. This can be performed with a dry cotton swab or small Calgiswab or by suctioning with a metal or plastic catheter. Cleansing by irrigation or flushing of the ear canal with a syringe or pulsatile irrigator (Water-Pik) may be more tolerable for some patients, but probably should be performed only when the tympanic membrane can be visualized due to the risk that the debris overlies an unseen perforation of the tympanic membrane.24

**Ear wick helpful.** After the ear canal has been debrided, consider the benefit of inserting an ear wick. Particularly if the ear canal is swollen, medication may not penetrate deep enough into the canal. In this situation, an ear wick, such as a Pope Oto-wick or Merocel XL (Medtronic Xomed, Inc, Jacksonville, Fla) hydrogel polymer, can enhance medication delivery directly to the entire ear canal.

The medication is absorbed into the ear wick, which delivers the drug to the infected skin (FIGURE). As swelling subsides, most ear wicks extrude spontaneously, usually within 12 to 36 hours.

**Common topical therapies**

Topical antibiotics commonly used for otitis externa are listed in the TABLE.

**Combination drugs containing aminoglycosides**

Combination topical drugs containing an aminoglycoside (neomycin), polymyxin B, and a corticosteroid were once the only antibiotic treatments for otitis externa approved by the US Food and Drug Administration.9 Neomycin is a bactericidal inhibitor of protein synthesis that is modestly active against *S aureus* but has minimal activity against *P aeruginosa.*25

In 1999, a prevalence study of aminoglycoside resistance in Europe found that 25% of the *S aureus* isolates tested were resistant to gentamicin.26 This study also found a high incidence (20%–38%) of gentamicin-resistant *P aeruginosa* isolates from Belgium, France, Italy, and Poland.26 Furthermore, in a prospective study of aural isolates of *P aeruginosa*, only 17.8% were susceptible to neomycin.27

<table>
<thead>
<tr>
<th><strong>TREATMENT (FORMULATION)</strong></th>
<th><strong>DOsing regimen</strong></th>
<th><strong>ADVANTAGES/DISADVANTAGES</strong></th>
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<tbody>
<tr>
<td>Neomycin sulfate/polymyxin B/ hydrocortisone (suspension/acidic)</td>
<td>4 times daily</td>
<td>Efficacious, extensively used; moderately expensive (AWP $65) Resistance and hypersensitivity increasing; potentially ototoxic; discourages compliance</td>
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<tr>
<td>Ofloxacin otic (solution/neutral)</td>
<td>Once or twice daily</td>
<td>Efficacious; no ototoxicity or hypersensitivity; favors compliance Resistance increasing; moderately expensive (AWP 5 cc $44.45; 10 cc $82.25)</td>
</tr>
<tr>
<td>Ciprofloxacin otic, with dexamethasone (suspension/acidic)</td>
<td>Twice daily</td>
<td>Efficacious; no ototoxicity with ciprofloxacin; favors compliance Resistance increasing; expensive (AWP $85)</td>
</tr>
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AWP, average wholesale price
Factors in choosing a topical antibiotic: evolving pathogen resistance, patient adherence, cost, possible hypersensitivity to neomycin

Polymyxin B is active against *P. aeruginosa* but not against *S. aureus*.28 Combining neomycin and polymyxin B broadens the spectrum of antimicrobial activity. Hydrocortisone, a low-potency steroid, purportedly helps reduce associated inflammation. Several ophthalmic preparations that contain a corticosteroid or an aminoglycoside, such as gentamicin or tobramycin, have been used off-label for otitis externa.3,20

Neomycin/polymyxin B/hydrocortisone has been shown in earlier clinical trials to be efficacious in up to 97% of cases of otitis externa.15 However, *S. aureus* has been increasingly resistant to aminoglycosides.30,31 More than 10% of *S. aureus* strains were resistant to gentamicin between 1987 and 1999.26,32

**Possible ototoxicity.** Aminoglycoside drops may cause vestibular ototoxicity when the tympanic membrane is perforated. Aminoglycosides instilled into the inner ear of test animals have been shown to damage cochlear inner and outer hair cells.33-35 Only a few similar case studies in humans have been published.36-38 Assessment of the tympanic membrane is often difficult because of excessive swelling of the external ear canal1 or patient noncooperation due to pain. In such cases—though tympanic membrane perforation is uncommon in otitis externa—physicians may want to avoid aminoglycosides.

**Beware hypersensitivity.** Drug hypersensitivity is another concern with topical aminoglycosides. Neomycin and thimerosal are among the antibiotic allergens most commonly detected on patch testing.39,40 Thimerosal is a preservative added to common otic formulations of neomycin/polymyxin B/hydrocortisone.41 In a retrospective review of patch testing reactions from 587 adult patients, neomycin sulfate and thimerosal elicited a hypersensitivity reaction in 53% and 18% of the patients, respectively.40

Hypersensitivity to these agents has also been shown in children by skin testing, although it is apparently rare clinically.42,43 A patch testing study involving 562 healthy schoolchildren found that 18.4% and 14.9% of the children had hypersensitivity on skin testing to neomycin sulfate and thimerosal, respectively.13 The high incidence of sensitization to neomycin and thimerosal may be related to the common use of neomycin as a topical ointment on skin abrasions and of thimerosal as a bacteriostatic preservative in immunizations in the past.

**Fluoroquinolones**

Oral fluoroquinolone antibiotics have been available for adults since 1990. The fluoroquinolones ofloxacin and ciprofloxacin were approved as topical therapy for otitis externa in 1997 and 1998, respectively. In addition, ofloxacin is indicated for otorrhea from the middle ear through an implanted tympanostomy tube (tube otorrhea).

Fluoroquinolones inhibit DNA gyrase and topoisomerase, which are required for bacterial DNA synthesis.23 These are broad-spectrum antibiotics that have good in vitro activity against both *S. aureus* and *P. aeruginosa*.23 An in vitro analysis of antimicrobial activity against clinical isolates of *S. aureus* and *P. aeruginosa* indicated that ofloxacin and ciprofloxacin were more active against these pathogens than was neomycin.44 The MIC90 values of ofloxacin and ciprofloxacin, respectively, were 1.0 µg/mL and 2.0 µg/mL for *S. aureus* and 2.0 µg/mL and 0.25 µg/mL for *P. aeruginosa*. In contrast, the MIC90 values of neomycin were 4.0 µg/mL for *S. aureus* and 16.0 µg/mL for *P. aeruginosa*. The MIC90 for polymyxin B against *P. aeruginosa* was 2.0 µg/mL.45

**Clinically equivalent to neomycin compound.** Recent clinical studies have shown that ofloxacin and ciprofloxacin are as efficacious as neomycin/polymyxin B/hydrocortisone in the treatment of otitis externa.21,22,46 In one study, the overall cure rate was 89% in patients treated with either ofloxacin (N=301) or neomycin/polymyxin B/hydrocortisone (N=300).21 In another study, ciprofloxacin was comparably effective as neomycin/polymyxin B/hydrocortisone therapy (93% [N=239] vs 87%
FAST TRACK

Remove debris from the ear canal, if not too painful, to maximize antibiotic contact.

**S aureus resistance has increased.** Fluoroquinolone use has been reported to lead to resistance in topical infections other than otitis externa. For example, the incidence of fluoroquinolone-resistant keratitis isolates of S aureus increased from 11% in 1990 to 28% in 1998. In another 5-year review of bacterial keratitis cases, the incidence of ciprofloxacin-resistant isolates of S aureus increased annually from 5.8% in 1993 to 35.0% in 1997, and the incidence of resistant isolates to ofloxacin increased from 4.7% to 35.0% over the same period. In contrast, P aeruginosa isolates remained susceptible to fluoroquinolones during the study periods. Resistance of P aeruginosa to fluoroquinolones has increased only slightly. Between 1991 and 1994, 0.44% of ocular isolates of P aeruginosa were resistant to ciprofloxacin, while 4.1% showed in vitro resistance between 1995 and 1998.

**Fluoroquinolone has advantages over combination therapy.** Ofloxacin and ciprofloxacin do not cause ototoxicity in humans and do not damage isolated cochlear cells, as can neomycin/polymyxin B/hydrocortisone combinations. In fact, both ofloxacin and ciprofloxacin/dexamethasone have been approved for use in patients with patent tympanostomy tube otorrhea.

Topical use of ofloxacin and ciprofloxacin is not associated with hypersensitivity, and hypersensitivity after the oral administration of these drugs is rare.

Finally, the once- or twice-daily dosage regimen of the topical fluoroquinolones markedly improves therapeutic adherence when compared with the 4-times-daily regimen of neomycin/polymyxin B/hydrocortisone.

**Preparations containing corticosteroids**

The benefit of corticosteroids for otitis externa is not well established, though they are added to many topical antibiotic preparations. In one study, the addition of hydrocortisone to neomycin and polymyxin-B reduced pain by approximately 1 day for patients with otitis externa. A combination of ciprofloxacin 0.3% and dexamethasone 0.1% administered twice daily is indicated for acute otitis externa, as was supported by an efficacy rate of 90.9% in a recent randomized, observer-masked, parallel-group, multicenter study in 468 children and adults. Although hypersensitivity to topical corticosteroids is well documented in published clinical studies, it seems to be rare in clinical practice.

**Special circumstances**

**Severe pain.** Inserting an ear wick and prescribing a topical fluoroquinolone plus an oral narcotic, such as acetaminophen with codeine, may be the optimal approach to resolving the infection and managing pain. Compared with neomycin combinations, ofloxacin and ciprofloxacin are somewhat more active in vitro against likely pathogens.

**Otitis externa with concomitant acute otitis media.** Antibiotics such as amoxicillin/clavulanate or cefdinir for acute otitis media should be used in addition to topical antibiotics.

**Otitis externa with mild localized cellulitis.** A standard oral antibiotic that covers staphylococci is recommended. Oral fluoroquinolones, for patients over 17 years of age, or intravenous ceftriaxone may also be considered for more severe cases associated with aural cellulitis.

For younger children, I have prescribed oral ciprofloxacin (off-label), which has been recently approved for children older than 12 months with complicated urinary tract infection. Outpatient parenteral ceftriaxone may alternatively be used for some cases of cellulitis if the patient is only moderately ill. If rates of community-acquired methicillin-resistant S aureus exceed 15% to 20% in the community, clinicians should consider empiric therapy initially with trimethoprim-sulfamethoxazole or clindamycin.

**Fungal infection.** If a patient develops (1) otitis externa refractory to 2 consecutive...
courses of topical antibiotics, or (2) exhibits a discharge that looks like a white, fluffy exudate, suspect a fungal infection and obtain a culture of the exudate. Empiric therapy with either oral fluconazole (Diflucan) or topical ciclopirox (Loprox) solution should be considered.

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The author has received grant/research support from Abbott Laboratories and GlaxoSmithKline, and has served on speakers’ bureaus for Abbott Laboratories.

**REFERENCES**


