UWTASP
tele-antimicrobial stewardship program

Pacific Northwest
Antibiotic
POCKET GUIDE

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Washington State Department of Health

UW Medicine

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The University of Washington Tele-Antimicrobial Stewardship Program (UW TASP) empowers antimicrobial stewardship teams throughout the Pacific Northwest by providing education, mentoring, community building, and resource sharing. By combining the resources available in our urban academic setting with the expertise of rural health providers, antimicrobial stewardship program implementation has been accelerated throughout the region with far reaching benefits to our community.

UW TASP has created the UW TASP Antibiotic Guide to provide prescribers with a tool to guide prescribing based on local, Pacific Northwest resistance-based data and expert opinion.

These guidelines are intended to support clinical decision-making but should not replace individual patient assessment or provider judgement. We encourage clinical discretion and welcome any feedback to improve these guidelines for future iterations.

For more information, please visit the UW TASP website at:
www.uwtasp.org

HOSPITAL USERNAME

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ANTIMICROBIAL STEWARDSHIP: GENERAL PRINCIPLES AND APPROACHES

The Agency for Healthcare Research and Quality (AHRQ) identifies 4 key moments in the decision making process to prescribe antimicrobials. This easy to remember approach can be used in most clinical settings and is outlined below.

CONSIDER THE FOUR MOMENTS OF ANTIBIOTIC DECISION MAKING

MOMENT 1: The Diagnosis

"Does this patient have an infection that requires antibiotics?"

Isolated changes in clinical status, lab values or vital signs ALONE should not trigger initiation of antibiotics. This is the time to pause and consider infectious and alternative non-infectious causes. Delirium in the elderly, aspiration pneumonitis, atelectasis, congestive heart failure, emboli, asymptomatic bacteriuria and/or pyuria are common examples of non-infectious conditions.

MOMENT 2: Initial Steps

"Have I ordered appropriate cultures before starting antibiotics?"

"What empirical antibiotic therapy should I initiate?"

"How do I ensure timely administration of appropriate empiric antibiotic therapy?"

Many community acquired infections can be treated empirically using local or regional guidelines tuned to surveillance microbiology data (i.e. antibiograms). Complicated, high-risk cases, recurrent infections, or patients at risk for drug resistant infections are most likely to benefit from reliable and timely microbiology. A standardized or institutional approach to treating common infections minimizes the delay to appropriate therapy.

NOTE *Procedures for optimal culture ordering, collection and reporting are detailed in the Guide to Utilization of the Microbiology Laboratory for Diagnosis of Infectious Diseases: 2018 Update by the Infectious Diseases Society of America and the American Society for Microbiology. The UW TASP guide does highlight some organism and syndrome-specific notes pertinent to empiric antimicrobial selection.*

MOMENT 3: Modification/De-escalation

"A day or more has passed. Can I stop antibiotics?"

"Can I narrow therapy?"

"Can I change from intravenous to oral therapy?"

Performing a regular antibiotic time-out for every patient on antibiotics with review of available microbiological data is the standard of care. Documentation in the medical record should include the anti-infective regimen, indication, the day of treatment, reasoning behind continuation or modification to regimen, plan for narrowing or transitioning to oral, and anticipated total duration. "The antibiotic time out" is best achieved through input by those involved in the prescribing, dispensing, administration and monitoring of antibiotics and hospital/clinic wide implementation. A team approach with comprehensive and clear documentation ensures the survival of the therapeutic plan through all transitions of care.

NOTE *Rapid diagnostics using molecular platforms and disease markers like procalcitonin have shortened the time from days to hours for usable lab/microbiology data. However, it is helpful to know your local lab tools and institutional protocols for result turn around and result interpretation.*

MOMENT 4: Duration

"What duration of antibiotic therapy is needed for this patient's diagnosis?"

Evidence supports shorter durations for common conditions. Most infections can be treated in 7 days or fewer. The total antibiotic duration count should include the first day appropriate empiric therapy was provided plus the days of targeted therapy. Minimizing excessive antibiotic exposure reduces the likelihood of drug side-effects, drug-drug interactions, antibiotic associated diarrhea including *C. difficile*, and resistance. Durations should be based on the current literature and initial clinical response.

NOTE *Because the majority of antibiotic prescribing in hospitals is for community-acquired pneumonia, ventilator-associated pneumonia, intra-abdominal infections, urinary tract infections, and cellulitis, these syndromes and their durations are specifically addressed in this guide. Duration updates are also highlighted in ambulatory conditions for upper and lower respiratory tract syndromes.*

ANTIBIOTIC RESISTANCE PEARLS

Regional resistance trends were utilized to drive agent selection for the UW TASP Antibiotic guide. Some customization of this guide may be warranted based upon your local antibiogram or drug formulary.

To learn more about regional antibiotic resistance, visit the Washington State Department of Health website for posted antibiograms.

https://www.doh.wa.gov/ForPublicHealthandHealthcareProviders/HealthcareProfessionsandFacilities/HealthcareAssociatedInfections/AntibioticResistance/Stewardship/Antibiograms

Following is a summary of observations for drugs and bugs that may help you in antibiotic selection:

GRAM-NEGATIVE BACTERIA AND ANTIMICROBIAL RESISTANCE:

Gram-negative bacteria like *E. coli, Klebsiella spp., Enterobacter, Acinetobacter,* and *Pseudomonas* are becoming increasingly drug resistant. Some of these organisms are intrinsically resistant due to structure or the production of specific beta-lactamases, but, over time, the repeated introduction of new genes on mobile plasmids is increasing the risk of unanticipated resistance profiles. This is especially concerning for empiric therapy for community-acquired infections since that therapy needs to cover essentially all probable bacterial pathogens. Although this is a very broad and complex topic, we are including a few examples below to help as you think through potential treatment options and/or interpret guidelines.

There has been a slow increase in resistance to the fluoroquinolone class of antimicrobials over the last 10-15 years. When considering treatment for empiric treatment options like complicated UTIs or gastrointestinal infections, fluoroquinolones may not be the best option. Surveillance data for Washington State demonstrates >15% fluoroquinolone resistance in *E. coli*, the most common community-acquired

Gram-negative pathogen. Treatment guidelines discourage empiric use of TMP/SMX for *E.coli* coverage when local susceptibility trends demonstrate resistance rates ≥20%, which is consistently observed in the Pacific NW. Similarly, ampicillin/sulbactam is no longer reliable for empiric coverage of *E. coli* due to rates of resistance commonly in the 30-40% range.

ESBL (extended spectrum beta-lactamase) Producers

Typical organisms: *E.coli, Klebsiella spp., Proteus*

Incidence: Accounts for at least 14% of E. coli in US hospitals according to a CDC report published in 2013

Resistance pattern: Can be susceptible to cephamycins (cefoxitin and cefotetan) and resistant to first and third generation cephalosporins

Recommended treatment: Although cephamycins show in-vitro susceptibility, they are **NOT** used for clinical ESBL infections. For serious infections due to ESBL-producing bacteria, carbapenems appear to be the best option, even if the organism is susceptible to drugs like piperacillin-tazobactam.

GRAM-POSITIVE BACTERIA AND ANTIMICROBIAL RESISTANCE:

Drug-resistance in Gram-positives is consistent with a longer history of resistance going back to at least the 1990s, most commonly in *Staphylococcus aureus* and *Enterococcus faecium*.

Staphylococcus aureus

S.a*ureus* are considered highly virulent organisms and can cause a variety of clinical syndromes from mild skin and soft tissue infections to life-threatening endovascular infections.

Methicillin-sensitive *Staphylococcus* aureus (MSSA)
Resistance pattern: *S. aureus* isolates sensitive to methicillin/oxacillin are also sensitive to nafcillin, ampicillin-sulbactam, amoxicillin-clavulanate, cefazolin and cephalexin. Often remains highly sensitive to the tetracycline class and TMP-SMX.

Recommended treatment: Nafcillin or cefazolin are appropriate first line therapies for treatment of serious MSSA infections. Although ceftriaxone is active against MSSA, it should not be used first line as clinical failures have been reported in the literature. The preferred oral agents for MSSA infections are cephalexin, dicloxacillin or TMP-SMX.

Methicillin-resistant *Staphylococcus* aureus (MRSA)
Resistance pattern: MRSA are resistant to essentially all beta-lactams, including cefazolin. Clindamycin susceptibility is variable and differences are often observed between MRSA isolates obtained in the hospital versus the community.

Recommended treatment: Vancomycin remains the drug of choice to treat hospitalized patients for MRSA infections. TMP-SMX maintains good susceptibility and is the preferred drug for ambulatory patients. The tetracycline class has remained relatively effective against MRSA, making doxycycline a reasonable choice for mild infections in patients with a sulfonamide allergy. Susceptibility results should be reviewed to ensure a negative D-test (indicating inducible clindamycin resistance) prior to utilization of clindamycin for definitive therapy for serious infections. Linezolid has reliable MRSA coverage but should be reserved for situations where intolerance or elevated MICs to vancomycin have been demonstrated or as an oral option for MRSA pneumonia. Extended duration of therapy (>10 days) with linezolid is cautioned due to increased risk of leukopenia, anemia, thrombocytopenia, lactic acidosis, and vision loss.

Rifampin should not be used as monotherapy for *S.aureus* due to rapid development of resistance and subsequent clinical failure. Rifampin may be an attractive option for *S.aureus* coverage in infections where biofilm production is concerning such as line sepsis or orthopedic post-operative infections with retained hardware, but only in combination with other anti-*S.aureus* agents.

Streptococci

Among *Streptococcus pneumoniae* in the United States, the reported rate of resistance to macrolides is 26.2%, resistance to TMP-SMX is 14.3%, clindamycin is 9.4%, and tetracycline is 16.2%. As a result, azithromycin and TMP-SMX are not recommended for empiric treatment options where coverage for *S. pneumoniae* is critical, such as most pediatric upper respiratory infections.

Groups A, B, C and G Streptococci are universally susceptible to penicillin and cefazolin; therefore, local testing and reporting is not necessary. *Streptococcus pyogenes* (Group A strep) and *S. agalactiae* (Group B strep) may exhibit inducible clindamycin resistance in up to 20% of cases. Confirm clindamycin susceptibility in these streptococcal infections prior to use.

Enterococcus

Enterococci are usually low virulence organisms and often over treated with antibiotics when isolated in non-sterile cultures. Urinary tract infections due to enterococci are often catheter or instrumentation-associated and bacteremia from a urinary source occurs infrequently. *Enterococcus* is a component of mixed flora in intra-abdominal and pelvic cultures and therapy specifically directed against this pathogen is generally not warranted. Non-antimicrobial treatments for enterococcal infections include catheter removal, percutaneous or surgical drainage, I&D and debridement.

Enterococcus faecalis

Resistance pattern: remain highly sensitive to ampicillin, nitrofurantoin, and vancomycin. Piperacillin, penicillin and amoxicillin activity can be extrapolated from ampicillin susceptibility. Note that trimethoprim-sulfamethoxazole (TMP-SMX) has unreliable activity against enterococci and is not tested due to the inherent ability of the organisms to take up exogenous folate.

Recommended treatment: cephalosporins and nafcillin **cannot** be used to treat enterococcal infections due to intrinsic resistance. The combination of ampicillin + gentamicin, ampicillin + ceftriaxone or vancomycin + gentamicin may be considered for endocarditis.

Enterococcus faecium

Resistance pattern: high-level beta-lactam resistance is common. Intrinsic resistance to cephalosporins and most carbapenems is the rule due to inner cell wall penicillin binding proteins (PBP). These organisms are often resistant to vancomycin as well, otherwise known as vancomycin-resistant enterococci (VRE).

Recommended treatment: Linezolid or daptomycin should be reserved for complicated VRE infections with or without bacteremia. Higher doses of daptomycin (10mg/kg) are recommended for severe enterococcal infections.

NOTE

- *Daptomycin should not be used to treat MRSA pneumonias due to drug degradation in the presence of surfactant.*
- *Ceftaroline, a newer cephalosporin with anti-MRSA activity, may be warranted in patients with persistent bacteremia. Cases with persistent bacteremia may benefit from expert ID consultation.*
- *Staphylococcus lugdenensis is a coagulase-negative staphylococcus similar to Staphylococcus epidermidis but has more invasive potential. It should not be treated as a contaminant until proven otherwise in clinical specimens. Most are treatable with oxacillin or cefazolin.*

EVALUATION AND DIAGNOSIS OF PENICILLIN ALLERGY FOR HEALTHCARE PROFESSIONALS

IS IT REALLY A PENICILLIN ALLERGY?

An accurate medication allergy history is the responsibility of every health care provider. It is imperative that antibiotic allergies be clarified, captured and, when appropriate, corrected in the electronic medical record. Formally, an allergy is a Type I immunoglobulin E-mediated adverse reaction that would be expected to be reproducible upon re-challenge.

A credible antibiotic allergy history includes two elements:

1. A specific recollection of the drug taken, the time elapsed between drug administration and drug reaction, and a physical description of the drug reaction,

   AND

2. Any signs and symptoms of a serious hypersensitivity reaction. The more specific and complete the symptoms of a drug reaction resembles an anaphylactic reaction, the more concerning and "credible" the history.

FACTS ABOUT PENICILLIN ALLERGY (TYPE 1, IMMUNOGLOBULIN E (IGE)-MEDIATED)

1. Approximately 10% of all U.S. patients report having an allergic reaction to a penicillin class antibiotic in their past.
2. However, many patients who report penicillin allergies do not have true IgE-mediated reactions. When evaluated, fewer than 1% of the population are truly allergic to penicillins.
3. Approximately 80% of patients with IgE-mediated penicillin allergy lose their sensitivity after 10 years.
4. Broad-spectrum antibiotics are often used as an alternative to penicillins. The use of broad-spectrum antibiotics in patients labeled "penicillin-allergic" is associated with adverse drug effects, sub-optimal antibiotic therapy, higher healthcare costs, and an increased risk for antibiotic resistance.
5. Correctly identifying those who are not truly penicillin-allergic can decrease unnecessary and inappropriate use of antimicrobials.

HISTORY AND PHYSICAL EXAMINATION

Before prescribing broad-spectrum antibiotics to a patient thought to be penicillin-allergic, evaluate the patient for true penicillin allergy (IgE-mediated) by conducting a history and physical.

Questions to ask to understand a patient's penicillin allergy:

- What medication were you taking when the reaction occurred?
- What kind of reaction occurred?
- How long ago did the reaction occur?
- How was the reaction managed?
- What was the outcome?
- Have you ever received amoxicillin, ampicillin or cephalexin since having the allergy?

Characteristics of an IgE-mediated (Type 1) reaction:

- Reactions that occur immediately or usually within one hour
- Hives: Multiple pink/red raised areas of skin that are intensely itchy
- Angioedema: Localized edema without hives affecting the abdomen, face, extremities, genitalia, oropharynx, or larynx
- Wheezing and shortness of breath

Anaphylaxis is a severe multisystem, IgE mediated reaction that occurs minutes to hours after exposure to an antigen exposure. Though several diagnostic criteria exist, the following are common involved systems:

Skin: Hives, flushing, itching, and/or angioedema

Respiratory: Cough, nasal congestion, shortness of breath, chest tightness, wheeze, sensation of throat closure or choking, and/or change in voice-quality (laryngeal edema)

Cardiovascular: Hypotension, faintness, tachycardia or less commonly bradycardia, chest pain, sense of impending doom, and/or loss of consciousness

Gastrointestinal: Nausea, vomiting, abdominal cramping, and diarrhea

ANTIBIOTIC CROSS REACTIVITY

Cross-reactivity refers to drugs with similar chemical structures that can induce similar allergic reactions. In the same manner that the allergy label has been misapplied to patients, cross-reactivity between chemically similar agents has been demonstrated via unreliable and imprecise diagnostic criteria, including reported allergies, which can overestimate the incidence of true drug allergies. Cross-reactivity requires confirmation with a drug-challenge. Reactions other than anaphylaxis, ex. delayed maculopapular skin eruption, are not cross-allergenic and do not create a contraindication for use.

NOTE *Patients with other severe hypersensitivity syndromes— like Stevens-Johnson syndrome, toxic epidermal necrolysis, serum sickness, acute interstitial nephritis, hemolytic anemia, and drug rash with eosinophilia and systemic symptoms (DRESS)—should* ***not*** *use the offending drug in the future. Skin tests and drug challenges are not appropriate for patients with these severe hypersensitivity syndromes.*

Cephalosporin use in penicillin-allergic patients - "What is the rate of cross-reactivity?"

Soon after the introduction of cephalosporins, anaphylaxis was reported in patients with prior reactions to penicillin. In the 1970s, a number of reviews examined the rate of allergic reactions to cephalosporins in penicillin allergic patients. One study found that 4.5% of about 16,000 patients exposed to penicillin had an allergy history; of the patients with allergy histories, 8% had a reaction to a cephalosporin. The 8% figure, rounded to 10%, has often been cited as the "rate" of cross-reactivity. A number of observations discredit the magnitude of this figure.

If a patient reports an allergy to penicillin, and a cephalosporin is ordered, the following recommendations can be made:

1. For a patient with a non-anaphylactic, non-IgE mediated penicillin reaction (a type II, III, IV or other reaction—hemolytic anemia, serum sickness, or maculopapular rash) cephalosporins can be given safely. This is especially true for a history of skin eruptions that do not involve itching or edematous wheals.
2. For patients with a history of a severe IgE-mediated penicillin reaction, the risk of a repeat reaction to an agent with a similar side chain is about 0.4%. With agents with dissimilar side chains the risk is nearly zero.

NOTE *Cefaclor, cefadroxil, and cephalexin are the cephalosporins that share the same side chain with ampicillin and therefore should* ***not*** *be given to patients with a history of anaphylaxis to penicillins. Cefotaxime, cefpodoxime, ceftriaxone and cefepime share the same side chains and have the potential for cross-reactivity amongst the third and fourth generations, but do not share with penicillin or amoxicillin.*

What about cross-reactivity among the penicillins, carbapenems or aztreonam?

Surprisingly little is known about the cross-reactivity between various penicillins (ampicillin, piperacillin, nafcillin, etc.). Amoxicillin, ampicillin, and penicillin have the most similar side chains. Until more information is available, a severe allergic reaction to penicillin is a contraindication to use of other penicillins.

Cross-reactivity between penicillin and carbapenems was initially controversial based upon inconsistent definitions and study design with imipenem. However, the most recent literature reports minimal risk. Specifically, with meropenem, data shows a risk of allergic reaction between 0%-0.9%. Patients

needing broad-spectrum beta-lactam therapy for high-risk severe infections or multi-drug resistant gram negative infections can safely receive meropenem even in patients with a definite history of anaphylaxis to penicillin.

Aztreonam is a monobactam with a low risk of cross-reactivity between other beta-lactam agents (with the exception of ceftazidime).

CONTENT SOURCE

Centers for Disease Control and Prevention, National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Division of Healthcare Quality Promotion (DHQP), https://www.cdc.gov/antibiotic-use/community/for-hcp/ Penicillin-Allergy.html; Page last reviewed: October 31, 2017 [Accessed March 2019]

Doherty K and Wilkerson T. Antibiotic Allergy and Cross Reactivity—A Review of the Literature. Jan 2013.

ORGAN SYSTEM:	SYNDROME:
Upper Respiratory	**Acute Otitis Media in Pediatrics**

SYMPTOMS AND/OR RISK FACTORS

Differential Diagnosis Details / non-AOM Conditions

- Middle ear effusion without inflammation suggests Otitis Media with Effusion (OME), a collection of non-infected fluid in the middle ear that may be due to viral URI, allergies, irritant exposure, eustachian tube dysfunction, or resolving AOM.
- Pain with mild traction to outer ear and normal appearing ear drum may indicate otitis externa. Inflammation of ear canal may be present but does not warrant systemic antibiotics.

AOM

- New onset otorrhea (not due to acute otitis externa)
- Mild bulging tympanic membrane and recent (less than 48 hours) onset of ear pain
- Moderate to severe bulging tympanic membrane
- Intense erythema of the tympanic membrane with presence of middle ear effusion
- Non-severe AOM is defined as mild otalgia for < 48 hours and temperature < 39°C (102°F)
- Severe AOM is defined as toxic-appearing child, moderate or severe otalgia, otalgia for > 48 hours, or temperature > 39°C (102°F) in past 48 hours
- Recurrent AOM (> 2 episodes in 6 months or > 3 episodes in 1 year) in children is an indication for referral for tympanostomy tube placement.

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

High-dose amoxicillin is recommended for pediatric otitis media because >10% *Strep pneumoniae* surveillance isolates are intermediate in Washington.

Culture of ear fluid is not typically indicated.

RECOMMENDED TREATMENT AND DURATION

The following cases should always be treated with antibiotics:

- AOM with otorrhea
- Severe AOM (unilateral or bilateral)
- Bilateral non-severe AOM without otorrhea in children 6-23 months
- Any AOM in infants < 6 months (infants < 2 months may require additional infectious work up)

FIRST LINE:

Amoxicillin (high-dose) 45 mg/kg PO BID (max 2000mg per dose)

NOTE: *For children with AOM and concurrent purulent conjunctivitis, use of amoxicillin in prior month, or history of recurrent treatment failures on amoxicillin, prescribe amoxicillin-clavulanate or a 2nd or 3rd generation cephalosporin.*

SECOND LINE:

Amoxicillin-clavulanate (ES 600mg/42.9mg) 45mg/kg PO BID (max 2000mg/dose)

Non-Type 1 β-Lactam Allergy: Cefprozil 15mg/kg PO BID (max 500mg/dose); Cefdinir 14mg/kg PO daily or 7mg/kg BID (max 600mg/day); Cefpodoxime 5mg/kg PO BID (max 200mg/dose); cefuroxime (Infants > 2 months) 15mg/kg PO BID (max 500mg/dose); Ceftriaxone 50mg/kg IM/IV daily (max 2gm/dose)

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

NOTE: *For children experiencing treatment failure (48-72 hours after initial antibiotic) alternatives include amoxicillin-clavulanate or ceftriaxone or clindamycin 10mg/kg PO TID (max 450mg/dose) or clindamycin PLUS 2nd or 3rd generation cephalosporin.*

DURATION:

- 1-3 days if treating with ceftriaxone IM/IV daily
- 5 days for non-severe AOM and age 2-5 years
- 7 days for non-severe AOM and > 6 years
- 10 days for severe AOM or age < 2 years

Consider watchful waiting without antibiotic therapy:

- For children > 23 months with either bilateral non-severe AOM without otorrhea or unilateral non-severe AOM without otorrhea.
- For children 6-23 months with unilateral non-severe AOM without otorrhea.

NOTE: *When watchful waiting is used, ensure follow-up and begin antibiotic therapy if patient is worsening or not improving within 48-72 hours*

SYMPTOMATIC TREATMENT for all patients:

- Extra rest, warm drinks, oral hydration
- Analgesics/antipyretics, as needed
  - Acetaminophen 15mg/kg PO q4-6hr PRN pain or fever, not to exceed 75mg/kg in 24 hours (max 3200 mg in 24 hours)
  - Ibuprofen 5-10mg/kg PO q8hr PRN pain or fever, not to exceed 30mg/kg in 24 hours (max 400mg/dose; 2400mg/day)
- Avoid cigarette smoke; offer smoking cessation resources to family members, if indicated

CONSIDERATIONS

- Ensure vaccinations are up to date.
- Cefuroxime oral suspension has been discontinued, consider cefprozil 15mg/kg PO BID (max dose 500mg) in children >6 months of age needing liquid antibiotic.
- Cefdinir, cefuroxime, cefpodoxime, cefprozil and ceftriaxone are highly unlikely to be associated with cross-reactivity with penicillin allergy on the basis of their distinct chemical structures.
- Consider ENT referral if no sign of improvement after 48-72 hours WITH failure of alternative agent.
- It is reasonable to treat AOM in adults with the same approach as pediatrics using adult dosing strategies for outlined regimens.

Best practices for communicating with patients

- Identify and validate patient's and parent's concerns.
- Provide clear recommendations including specific symptom treatment and contingency plan for if symptoms worsen.
- Confirm agreement and answer questions.
- Provide education about antibiotic use and associated risks, including bacterial resistance, and *C. difficile.*
- Visit CDC's Common Illnesses index at *https://go.usa.gov/xRPXH* for patient education materials.

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(adopted from Washington State Department of Health guideline DOH 420-197 Aug 2017)

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2. *Limb CJ, et al., Acute otitis media in adults. In: UpToDate, Libman H (Ed), UpToDate, Waltham, MA. Accessed on February 16, 2017.*

ORGAN SYSTEM:	SYNDROME:
Upper Respiratory	**Sinusitis in Pediatrics and Adults**

SYMPTOMS AND/OR RISK FACTORS

Cardinal Criteria for Bacterial Sinusitis

Must have purulent nasal discharge

PLUS

Nasal obstruction AND/OR facial pain/pressure/fullness

AND

Persistent & not improving (>10 days) OR symptoms worsen within 10 days after initial improvement (double worsening from a typical upper respiratory infection that lasted 5-6 days)

NOTE: *thick, colored, or purulent nasal secretions do NOT necessarily indicate bacterial infection*

Items to consider for Risk of Antibiotic Resistance:

- Prior Abx in past 30 days
- Age <2 or >65
- Comorbidities
- Prior hospitalization in past 5 days
- Attend daycare
- Immunocompromised
- Moderate to severe or prolonged signs and symptoms
- Failure of prior ABX treatment
- Frontal or sphenoidal sinusitis

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

- > 95% of cases are of viral origin and do not warrant antibiotics.
- Approximately ¼ of *H. influenza* isolates produce beta-lactamases and are resistant to amoxicillin.
- > 10% of *Strep pneumoniae* surveillance isolates are non-susceptible to standard dosing of amoxicillin warranting higher dose in select patients.
- Macrolides are NOT recommended for empiric therapy due to high rates of resistance among *S. pneumoniae.*
- Sulfamethoxazole/Trimethoprim is NOT recommended for empiric therapy due to high rates of resistance to *S. pneumoniae* and *H. influenzae.*
- Routine coverage for MRSA is NOT recommended for initial empiric therapy.

NOTE: *Endoscopic-guided culture and/or empiric Staphylococcus aureus (bactrim or doxycycline) should be considered in patients who have had recent sinus surgery.*

RECOMMENDED TREATMENT AND DURATION

Watchful waiting:

- Acceptable to observe mild bacterial sinusitis for 7 additional days before prescribing antibiotic if follow up is assured and focus instead on symptomatic treatment.
- Consider delaying the initiation of ABX for any severity of symptoms.
- Initiate tx if condition fails to improve by 3 days in children or 7 days in adults.
- Consider wait-and-see-prescription (WASP).

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

Exceptions to watchful waiting:

- Patients with Chronic Rhinosinusitis or recurrent Acute Rhinosinusitis in multiple chronic conditions such as: asthma, ciliary dyskinesia, cystic fibrosis, or immunocompromised state.
- Watchful waiting may not be reasonable for advanced age, impaired cardiopulmonary status or multiple co-morbidities and overall poor general health.

If cardinal criteria are met and at least 10 days of symptoms or double worsening occurs:

FIRST LINE ADULT:

Amoxicillin-clavulanate 875mg/125mg PO BID x 5 days

SECOND LINE ADULT:

β-Lactam Allergy: Doxycycline 100mg PO BID; or Clindamycin 300mg PO TID plus Cefpodoxime 200mg PO BID x 5 days; or Levofloxacin 500mg PO Q 24 Hours x 5 days

NOTE: *if cefpodoxime unavailable, substitute alternative 2nd or 3rd generation cephalosporin in above clindamycin combination regimen.*

At Risk for Antibiotic resistance: Amoxicillin-clavulanate 2gm PO BID; if high-dose extended release formulation not available: Amoxicillin-clavulanate 875mg/125mg PO BID plus Amoxicillin 1gm PO BID x 5 days; or Levofloxacin 500mg PO Q 24 Hours

UPDATE: *Fluoroquinolone FDA Safety Alert: Disabling & potentially permanent adverse effects outweigh benefit in sinusitis. Only use levofloxacin when no other alternatives exist.*

FIRST LINE PEDIATRIC:

Amoxicillin/clavulanate: 22.5 mg/kg PO BID x 10 days

SECOND LINE PEDIATRIC:

Non-Type 1 β-Lactam Allergy: Clindamycin 10mg/kg PO TID plus Cefdinir 14mg/kg/day x 10 days; or Levofloxacin [max dose of 500mg] 6 months to 5 years old: 8-10mg/kg PO BID x 10 days or 5 to 16 years of age: 8-10mg/kg PO Q 24 Hours x 10 days

At Risk for Antibiotic resistance: Amoxicillin-clavulanate (High dose-ES 600mg/42.5mg/5mL) 45mg/kg PO BID x 10 days or use same regimen options for Non-Type 1 β-Lactam Allergy option in pediatrics above.

Symptomatic Relief/ Adjunctive Treatment:

- Intranasal saline irrigation is safe and effective for symptom relief & does not lead to resistance.
- Intranasal corticosteroids are recommended for patients with h/o allergic rhinitis at standard approved dosing strategies.
- Control pain/fever with ibuprofen or acetaminophen.
- Nasal decongestants like oxymetazoline 1-3 sprays each nostril daily for up to 1 week if used concomitantly with intranasal steroids are safe and effective in adults with sinusitis.

CONSIDERATIONS

Identify and validate patient's concerns and provide clear recommendations including specific symptom treatment and contingency plan for if symptoms worsen.

During follow-up, if patient worsens or lack of improvement at 7 days from presentation:

- Reassess and confirm diagnosis, exclude other causes, and detect complications
- If watch and wait management, initiate FIRST LINE treatment
- If FIRST LINE treatment already completed, consider treatment from "At risk for ABX resistance"

During follow-up, if NO improvement after 2 courses of antibiotics or if concern for orbital/CNS complications of bacterial sinusitis, order contrast-enhanced CT scan (preferred) or MRI of the paranasal sinuses and refer to the appropriate specialist.

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(adopted and updated from WS DOH 420-194 Nov 2017)

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ORGAN SYSTEM:	SYNDROME:
Upper Respiratory	**Pharyngitis in Pediatrics & Adults**

SYMPTOMS AND/OR RISK FACTORS

Symptoms

- Abrupt onset of sore throat
- Headache
- Myalgia
- Occasionally nausea/vomiting/abdominal pain followed by spontaneous resolution in 2-5 days

Physical Exam consistent with **Bacterial** Pharyngitis

- Patchy tonsillopharyngeal exudate
- Anterior cervical adenitis (tender nodes)
- Tonsillopharyngeal inflammation
- Fever >100.4 F
- Palatal Petechia
- Scarlatiniform rash
- Absence of cough

NOTE: *If severe signs/symptoms (drooling, dysphonia, "potato" voice, neck swelling) consider: epiglottitis, peritonsillar abscess, retropharyngeal abscess, submandibular space infections, or primary HIV. Obtain lateral neck x-ray, and consider transfer to the emergency department.*

Viral Features

- Conjunctivitis
- Rhinorrhea
- Coryza
- Cough
- Oral ulcers
- Hoarseness (laryngitis)
- Viral exanthema
- Diarrhea
- Ear pain

NOTE: *> 95% of pharyngitis cases are of viral etiology and do not require antibiotics. Provide symptomatic relief.*

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Test:

- Testing for Group A Streptococcus (GAS) is NOT recommended for acute pharyngitis with clinical & epidemiologic features that strongly suggest a VIRAL etiology.
- Routine use of back up throat cultures for those with a negative RADT is NOT necessary for adults; there is a low incidence of GAS pharyngitis in adults & risk of subsequent acute rheumatic fever is exceptionally low.
- Rapid Diagnostic Test (RADT) Recommended for adults with two or more symptoms and for children with signs and symptoms of strep throat who do not have viral symptoms.
- Reflex/Back up throat culture for negative RADT is only indicated in children/adolescence (3-15 years), patients at high-risk for severe disease (eg. poorly controlled diabetes, immunocompromised, on chronic corticosteroids), or those in close contact with elderly, infants or immunocompromised individuals.

NOTE: *It is NOT recommended to test for GAS under the age of 3 years.*

RECOMMENDED TREATMENT AND DURATION

NOTE: *treat patients who are RADT or throat culture positive or those with known exposure 2 weeks prior to symptom onset.*

FIRST LINE PEDIATRIC:

- Pen VK 250mg PO BID - TID (>27kg 500mg BID - TID) x 10 days
- Amoxicillin 50mg/kg PO daily or divided in 2 doses (max 1gm/day) x 10 days
- Penicillin G Benzathine (<27kg) single IM dose 600,000 units x 1 dose

  UPDATE: *Drug shortage of IM Penicillin G Benzathine warrants oral treatment options as first line consideration.*

SECOND LINE PEDIATRIC:

- Non-Type 1 β-Lactam Allergy: Cephalexin 20mg/kg PO BID (max 500mg/dose) x 10 days
- Type 1 β-Lactam Allergy: Azithromycin (2-15 years of age) 12mg/kg PO once, then 6mg/kg PO daily days 2-5 (max 500mg/dose); Azithromycin 20mg/kg PO once daily (max 1000mg/dose) x 3 days; or Clindamycin 7mg/kg PO TID (max 300mg/dose) x 10 days

FIRST LINE ADULT:

- Pen VK 500mg PO BID-TID x 10 days
- Amoxicillin 1000mg PO daily OR 500mg PO BID x 10 days
- Penicillin G Benzathine (>27kg) 1.2 million units IM x 1 dose

SECOND LINE ADULT:

- Non-Type 1 β-Lactam Allergy: Cephalexin 500mg PO BID x 10 days
- Type 1 β-Lactam Allergy: Azithromycin 500mg PO on day one, 250mg PO daily on days 2-5; Azithromycin 500mg PO daily x 3 days, or Clindamycin 300mg PO TID x 10 days

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

Symptomatic Relief for all Patients (viral or bacterial infections):

- Rest
- Adequate fluid intake
- Antipyretics (no ASA under age 2)
- Magic mouthwash
- > 6yrs of age: gargle with warm salt water
- > 3yrs of age: sucking on hard candy

NOTE: *Medicated throat lozenges/sprays (not recommended in children/adolescents)*

CONSIDERATIONS

- Individual will be contagious for 24 hours after starting antibiotic tx.
- Treatment for non-symptomatic GAS carriers is NOT routinely recommended.
- Testing or empiric tx of asymptomatic household contacts is NOT routinely recommended.
- There is no evidence of benefit for glucocorticoids in children or adolescents. Short term dosing may be beneficial in adults.
- Treatment for Group C & G are the same recommendations.
- Best Practices for Communicating with Patients:
  - Identify and validate patient's and parent's concerns.
  - Provide clear recommendations including specific symptom treatment and contingency plan for if symptoms worsen.
  - Confirm agreement and answer questions.
  - Provide education about antibiotic use and associated risks, including bacterial resistance and *C. difficile.*

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(adopted from Washington State Department of Health DOH 420-198 Aug 2017)

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ORGAN SYSTEM:	SYNDROME:
Lower Respiratory	**Acute Uncomplicated Bronchitis in Adults**

SYMPTOMS AND/OR RISK FACTORS

Presenting Symptoms:

- Cough > 5 days in a patient WITHOUT COPD
- Purulent sputum occurs in 50% of cases and does NOT necessarily indicate bacterial infection
- Low-grade fever is common early in illness (<100.5 F or <38C)
- Diffuse wheezes or rhonchi on exam, but NOT rales or signs of consolidation
- Mild dyspnea
- Chest wall pain due to coughing

Comorbidities to consider:

- COPD
- Asthma
- Elderly (> 75 years)
- Immunocompromised
- Heart failure

Testing:

- Vital signs including SpO2
- Obtain CXR if: hemoptysis, ill-appearing, focal abnormality on auscultation, age >70, RR >24 bpm, temperature > 100.4F or >38C for longer than 4 days or recurrent after having resolved for longer than 24 hours, HR > 100 bpm, abnormal oxygen saturation, cough not improving after > 6-8 weeks
- Consider smoking history when considering CXR
- Procalcitonin (if available) may help confirm decision to withhold antibiotics

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Bronchitis is a self-limited inflammation of the bronchi due to respiratory infection by viruses (>90% of cases)

- Influenza A or B
- Parainfluenza
- Human metapneumovirus
- Rhinovirus
- RSV
- Pertussis
- Consider influenza PCR during flu season if high risk or <48 hours of symptoms
- Consider pertussis PCR if paroxysms, post-tussive emesis, inspiratory whoop or known exposure to pertussis case. Report suspect, probable or confirmed pertussis to local public health.
- Respiratory pathogen testing is discouraged in uncomplicated acute bronchitis

NOTE: *the most common causes of acute uncomplicated bronchitis DO NOT require antibiotics*

RECOMMENDED TREATMENT AND DURATION

NOTE: *Antibiotic therapy may be indicated for bronchitis in patients with comorbidities such as immunosuppression, COPD/chronic bronchitis, cystic fibrosis, or other underlying lung disease other than asthma. Recommendations for these patients is beyond the scope of this guideline.*

Symptoms without comorbidities present < 14-21 days:

- Guaifenesin 100mg/5ml PO Q4H prn cough
- Dextromethorphan 10-20 mg Q4H prn cough (max 120 mg/24hr)

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

UPDATE: *Narcotic medications should not be used for cough suppression in acute bronchitis*

- Albuterol inhaler 90 mcg/inhalation 1-2 puffs PO QID prn difficulty breathing and wheezes present on exam in patients with asthma or underlying pulmonary disease

Symptoms and comorbidities present:

- Evaluate for pneumonia or COPD exacerbation or alternative causes
- If positive evaluation, treat accordingly
- If negative evaluation, follow guideline for symptoms without comorbidities

Adjunctive medications:

- Ibuprofen 400mg PO Q6-8H prn pain or inflammation
- Naproxen 500mg PO Q12H prn pain or inflammation
- Acetaminophen 325mg-650mg PO Q6h prn pain

CONSIDERATIONS

- Expected duration of cough is 2-3 weeks (average 18 days). In addition to cough due to acute bronchitis, persistent cough, especially cough lasting > 6-8 weeks, may be a sign of another disease process ranging from minor to serious, such as post-nasal drip syndrome, medication use (e.g., lisinopril), irritant exposure, asthma, Gastroesophageal Reflux Disease (GERD), smoking or second-hand smoke exposure, chronic bronchitis, bronchiectasis, or malignancy.
- Antihistamines are NOT effective for bronchitis.
- Provide patient education on rationale for not prescribing antibiotics, expected duration of symptoms, importance of smoking cessation and smoke-free environment, avoidance of irritants, adequate hydration, rest, humidified air, and to follow-up for worsening symptoms. Describe the diagnosis as "viral illness" or "chest cold".

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ORGAN SYSTEM:	SYNDROME:
Lower Respiratory	**Pneumonia in Pediatrics**

SYMPTOMS AND/OR RISK FACTORS

Initial Testing/Imaging

- Vital Signs: Temp, BP and Pulse Oximetry

NOTE: *No routine labs or CXR are indicated for children well enough to be managed outpatient.*

- Labs:
  - Blood work: CBC with differential, CRP, blood cultures if not fully immunized OR fails to improve after initiation of antibiotics
  - Viral Testing: Influenza PCR during influenza season
  - If atypical pathogen suspected: PCR Respiratory Panel if available
  - Sputum gram stain and culture: if intubating, collect at time of initial ET tube placement; consider testing in older children who can produce sputum sample
  - Urinary antigen detection testing is not recommended in children; false-positive tests are common.
- Radiography:
  - AP and lateral CXR if failure to improve on initial antibiotic therapy
  - AP and lateral CXR 4-6 weeks after diagnosis if recurrent pneumonia involving the same lobe

Criteria for Outpatient Management

- Mild CAP: no signs of respiratory distress and SpO2 ≥=90% on room air
- Able to tolerate PO
- No concerns for pathogen with increased virulence (ex. CA-MRSA)
- Family able to carefully observe child at home, comply with therapy plan, and attend follow up appointments

Inpatient Admission Criteria

PEDIATRIC FLOOR

- Respiratory distress
- SpO2 <90% on room air
- Unable to tolerate PO
- Suspected or documented CAP caused by pathogen with increased virulence (ex. CA-MRSA)
- Concerns about observation at home, inability to comply with therapy, inability to be followed up

PICU

- Respiratory support: Intubated or requiring non-invasive positive pressure ventilation
- Concern for respiratory failure
- Concern for sepsis
- FiO2 needs HFNC >50% to keep saturation ≥92%
- Altered mental status

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

- The most common suspected pathogens for bacterial pneumonia include: *Streptococcus pneumonia* and *Haemophilus influenzae.*
- For suspected atypical pneumonia, the most common organisms include: *Mycoplasma pneumoniae* or *Chlamydophila pneumoniae*. For children ≥ 5 years of age, empirically add a macrolide if atypical pneumonia cannot be ruled out.
- For suspected viral pneumonia, the most common pathogens include: Influenza A & B, Adenovirus, Respiratory Syncytial Virus (RSV) and Parainfluenza.

RECOMMENDED TREATMENT AND DURATION

UNCOMPLICATED PNEUMONIA

Previously healthy and fully immunized children:

Inpatient Treatment:

FIRST LINE:

Ampicillin 50mg/kg IV q6hr (max 12g/day)

SECOND LINE:

- Non-Type 1 β-Lactam Allergy: Ceftriaxone 50mg/kg IV q24hr (max 2g/day)
- Type 1 β-Lactam Allergy: Levofloxacin <5 years: 10mg/kg IV BID (max dose 750mg/day); >5 years: 10mg/kg IV q24hr (max dose 750mg/day)

Outpatient Treatment:

FIRST LINE:

Amoxicillin 45mg/kg PO BID (max dose 4000mg/day)

SECOND LINE:

- Non-Type 1 β-Lactam Allergy: Cefuroxime 15mg/kg PO BID (max 1000mg/day) or cefprozil 15mg/kg PO BID (max dose 500mg) in children > 6 months of age needing a liquid formulation
- Type 1 β-Lactam Allergy: Levofloxacin <5 years: 10mg/kg PO BID (max dose 750mg/day); >5 years: 10mg/kg PO daily (max dose 750mg/day)

COMPLICATED PNEUMONIA

Not appropriately immunized with PCV13 & Hib or there is a suspicion for *H. influenzae* or severe disease and/or complicated pneumonia:

Inpatient Treatment:

FIRST LINE:

Ceftriaxone 50mg/kg IV q24hr (max 2g/day)

SECOND LINE:

Type 1 β-Lactam Allergy: Levofloxacin <5 years: 10mg/kg IV BID (max dose 750mg/day); >5 years: 10mg/kg IV q24hr (max dose 750mg/day)

Outpatient Treatment:

FIRST LINE:

Amoxicillin/clavulanate <40kg: (ES 600mg/42.5mg/5mL) 45mg/kg PO BID or 15mg/kg PO TID (max dose 4000mg/day); >40kg: 875mg/125mg PO BID PLUS Amoxicillin 1g PO BID

SECOND LINE:

Same as second line oral options recommended for outpatient treatment noted above as cephalosporin or fluoroquinolone adequately covers pneumococcus and *H. influenzae.*

- Uncomplicated pneumonia: complete a 10 day course of FIRST LINE therapy. Although 10 day durations have been best studied in children, shorter courses (5-7 days) may be considered for mild disease and in those managed as outpatient.
- Complicated pneumonia: duration is dependent on clinical response, in general 2-4 week course. For suspicion of Staphylococcus aureus:
- In addition to one of the above antibiotics, ADD: Clindamycin 10mg/kg IV q6hr (max 900mg/dose)
- For PICU or Severe Infection, ADD Vancomycin 15mg/kg IV q6hr (max 4g/day)

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

For pneumonia in children > 5 years of age and atypical pneumonia cannot be ruled out:

FIRST LINE:

ADD azithromycin 10mg/kg IV/PO daily for 1-2 days then transition to oral step down if possible (max 500mg/dose). A 3-day total azithromycin course is sufficient for atypical coverage.

SECOND LINE:

(For children > 7 years only) ADD Doxycycline 1-2 mg/kg PO BID (max dose 200mg/day) for 7-10 days.

CONSIDERATIONS

- Viral pneumonia is most common in children < 5 years of age. Antibiotics are not typically necessary. If influenza positive, treat with oseltamivir.
- Children should show clinical signs of improvement within 48-72 hours allowing de-escalation of therapy based on available culture results and consideration of transition to oral step-down therapy.
- If no improvement or worsening, pursue further diagnostic work up as indicated. Consider broadening antibiotics and formal infectious disease consultation.

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ORGAN SYSTEM:	SYNDROME:
Lower Respiratory	**Pneumonia in Adults**

SYMPTOMS AND/OR RISK FACTORS

Symptoms: productive cough, chest pain, dyspnea (SOB), diminished breath sounds, crackles not cleared with coughing, abdominal pain, with or without fever

Assess: Chest X-ray; pulse oximetry

Adult **CURB-65** Score (0-1) Manage as Outpatient:

Confusion

Blood **U**rea nitrogen > 20 mg/dL

Respiratory rate ≥ 30 breaths/min

Blood pressure SBP < 90 or DBP ≤ 60 mmHg

Age ≥ **65** years

HIGH-RISK FOR MDRO PNEUMONIA

Receipt of IV antibiotics in preceding 90 days has been defined as a sole risk factor for multi-drug resistant organisms (MRDO) by the most recent IDSA clinical guidelines for nosocomial pneumonia.

Consider prior fluoroquinolone exposure in previous 90 days when selecting empiric drug regimen; consider avoidance of repeated fluoroquinolone exposure. Provide additional gram negative coverage for those with risk factors for MDRO's.

Consider the Drug Resistance in Pneumonia (DRIP) Score for predicting risk of community-acquired pneumonia due to drug-resistant pathogens. Scoring tool can be found on-line at *www.mdcalc.com/drug-resistance-pneumonia-drip-score.*

NOTE: *Compared to HCAP, the DRIP Score is more sensitive (82% vs 79%), more specific (81% vs 65%), and decreases use of unnecessary extended-spectrum antibiotics by 46%.*

Major Risk Factors (each counts for 2 points):

- Antibiotic use within 60 days
- Long-term care resident (not including assisted living or group home facilities)
- Tube feeding (NG, NJ or PEG)
- Prior drug-resistant pneumonia diagnosis within 1 year

Minor Risk Factors (each counts for 1 point):

- Hospitalization within 60 days
- Chronic pulmonary disease
- Poor functional status (Karnofsky Performance Status < 70 or non-ambulatory status)
- H2 Blocker or PPI within 14 days
- Active wound care at time of admission
- MRSA colonization within 1 year

Management based upon total score:

- A patient with a DRIP Score <4 can effectively be treated without broad-spectrum antibiotic coverage.
- A patient with a DRIP Score of ≥4 is more likely to require broad-spectrum antibiotic coverage.

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Send sputum for gram stain & culture, CXR, urinary pneumococcal antigen, urinary legionella antigen, and blood cultures for hospitalized patients.

Send nasal swab for rapid influenza testing.

Respiratory PCR and/or procalcitonin (PCT) may be helpful if unclear diagnosis of pneumonia or acute exacerbation of COPD.

Most Common Etiologies:

Bacterial: *S. pneumonia*, Mycoplasma, *H. influenza, Chlamydophila pneumoniae*

Respiratory viruses: Influenza A & B, adenovirus, respiratory syncytial virus, parainfluenza

Structural lung disease such as bronchiectasis or exacerbations of COPD with multiple courses of antibiotics and/or chronic steroid use may warrant coverage for *Pseudomonas aeruginosa*

RECOMMENDED TREATMENT AND DURATION

Community-acquired pneumonia (non-aspiration risk)

FIRST LINE:

- Ceftriaxone 1gm IV q24hr x 5 days PLUS Azithromycin 500mg PO/IV q24hrs x 3 days
- Amoxicillin 1gm PO TID PLUS Azithromycin 500 mg PO daily preferred for ambulatory patients or as an oral option de-escalation approach to complete IV inpatient therapy.

SECOND LINE (beta-lactam allergy):

- Levofloxacin 750 mg PO/IV q24hrs x 5 days.
- Cefuroxime 500mg PO BID or cefpodoxime 200mg PO BID as the alternative for ceftriaxone or amoxicillin is preferred for ambulatory patients or as an oral option de-escalation approach to complete IV inpatient therapy.

NOTE: *In previously healthy individuals with no recent antibiotic therapy within previous 3 months, and no risk for drug-resistant S. pneumoniae, Doxycycline 100mg PO BID monotherapy may be considered.*

UPDATE: *Azithromycin is provided for atypical coverage and should not be relied upon as monotherapy for ambulatory or inpatient management of pneumonia due to increasing Streptococcus pneumoniae resistance.*

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

Community-acquired pneumonia (non-aspiration risk with risk factors for MDROs)

FIRST LINE:

Cefepime 2gm IV q8hr (1gm q8hr EI) AND Azithromycin 500mg PO/IV q24hr

SECOND LINE:

Levofloxacin 750mg PO/IV q24hr +/- Aztreonam 2gm IV q8hr

Ventilator Associated Pneumonia

- Early onset: (≤ 4 days of hospitalization or ventilation): Ceftriaxone 1gm IV q24hr
- Late onset: (> 4 days of inpatient stay)
  Treat as High-risk for MDRO's: Cefepime 2gm IV q8hr +/- Vancomycin/Linezolid if history of MRSA infection/colonization

Typical duration for VAP regardless of onset is 7 days

Chronic Aspiration Pneumonia

NOTE: *Prophylactic antibiotics do not offer any clinical benefit for patients with acute aspiration pneumonitis.*

Aspiration Pleuropulmonary Syndrome (Anaerobic coverage is clearly indicated only in the classic aspiration pleuropulmonary syndrome in pts with a h/o LOC as a result of EtOH/drug overdose or after seizures in pts with concomitant gingival disease or esophageal motility disorders)

FIRST LINE:

Ampicillin/sulbactam 3gm IV q6hr for 7-21 days

SECOND LINE:

- Non-severe penicillin allergy: Ceftriaxone 1gm IV q24hr AND metronidazole 500mg PO/IV q8hr
- Severe penicillin allergy: Moxifloxacin 400mg PO/IV q24hr

NOTE: *If history of MRSA infection/colonization, consider adding vancomycin or linezolid. If MDRO risk factors: Piperacillin/tazobactam 4.5gm IV q6hr (3.375gm IV q8hr xtended infusion) as alternative to ampicillin/sulbactam is appropriate.*

CONSIDERATIONS

- During flu seasons, send Flu testing and then give oseltamivir 75mg PO/NG BID. Higher doses of oseltamivir (ie. 150mg BID) in critically ill or obese patients have not been associated with improved outcomes.
- Yeast in sputum rarely represents true infection.
- If MRSA nares swab or sputum is negative for MRSA, discontinue vancomycin.
- Anaerobic coverage such as piperacillin-tazobactam is not usually necessary for CAP, HAP or VAP.
- Narrow therapy based on microbiology.
- Consider formal ID consultation if not clinically improving.
- Regardless of type of pneumonia, consider MRSA coverage if post-influenza pneumonia (days to weeks) and necrotizing/life-threatening presentation. Ensure regimen targets *S. pneumoniae* and *H. influnezae* as well.
- CF or Lung Transplant patients likely require pulmonary transplant or transplant infectious diseases consult.

ANTIBIOTIC DOSING CONSIDERATIONS

If evidence of only pneumococcal infection (including bacteremia), de-escalate to amoxicillin 1 gm PO TID and treat for total appropriate abx duration of 5-7 days.

NOTE: *High dose amoxicillin is appropriate for Streptococcus pneumoniae, cefinase negative H. influenzae or M. catarrhalis. If CAP complicated by empyema, asplenia or elevated Streptococcus pneumoniae MIC 2-4, add amoxicillin 1gm PO BID PLUS amoxicillin/clav 875mg PO BID.*

If MRSA coverage is warranted, base initial vancomycin dosing on patient's actual body weight using the following weight cutoff values.

Vancomycin Loading Dose:
<50 kg: Vancomycin 1gm IV x 1
50-70 kg: Vancomycin 1.5gm IV x 1
>70 kg: Vancomycin 2gm IV x 1

Vancomycin maintenance dosing should be established in collaboration with clinical pharmacy services.

DURATION

Duration of therapy is typically dependent upon initial clinical improvement and underling health with or without structural lung disease.

- 5 days for patients without immunosuppression or structural lung disease
- 7 days for patients with moderate immunosuppression or structural lung disease
- 10-14 days for poor clinical response, initial inappropriate tx, or significant immunosuppression

NOTE: *Patients should be afebrile for 48-72hr and demonstrate signs of clinical stability before therapy is discontinued.*

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ORGAN SYSTEM:	SYNDROME:
Intra-abdominal	**Intra-abdominal Infections in Adults Inpatient**

SYMPTOMS AND/OR RISK FACTORS

High Risk/Severe Criteria

Albumin <2.5

Age >70 years

Immunocompromised state

Severe sepsis/septic shock

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Intra-abdominal infections are usually of a polymicrobial process and may include the following pathogens:

- Enterobacteriaceae
- Enterococcus sp.
- Anaerobes (including Bacteroides sp.)

NOTE: *Anaerobes are less significant for biliary sources unless bile duct to bowel anastomosis or fistula present*

Routine cultures are not recommended for community-acquired infections.

Cultures SHOULD be obtained in patients with nosocomial infection or who require operation for prior treatment failure.

RECOMMENDED TREATMENT AND DURATION

EXTRA-BILIARY SOURCE:

appendicitis, diverticulitis, bowel perforation with peritonitis

Extra-biliary Source MILD-MODERATE Risk

FIRST LINE

Ceftriaxone 2gm IV q24hr PLUS Metronidazole 500mg IV q8hr

SECOND LINE

(Type 1 β-Lactam Allergy): Levofloxacin 500mg IV q24hr PLUS Metronidazole 500mg IV q8hr

Extra-biliary Source HIGH RISK/SEVERE

FIRST LINE

Piperacillin-tazobactam 4.5gm IV q6hr (or extended infusion)

SECOND LINE

(Type 1 β-Lactam Allergy): Levofloxacin 500mg IV q24hr PLUS Metronidazole 500mg IV q8hr +/- Aztreonam 2mg IV q8hr

NOTE: *IF previous colonization or concerns for highly resistant GNRs, consider meropenem 1gm IV q8hr as a substitute for piperacillin-tazobactam or additional GNR coverage to levofloxacin.*

Duration of therapy

4 days with adequate surgical source control

5 days for uncomplicated diverticulitis

If retained focus of infection, duration should be guided by clinical response.

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

BILIARY SOURCE:

cholecystitis, cholangitis

Biliary source MILD-MODERATE Risk

FIRST LINE

Ceftriaxone 2gm IV q24hr

SECOND LINE

(Type 1 β-Lactam Allergy): Levofloxacin 500mg IV q24hr

NOTE: *If bilio-enteric anastomosis present add metronidazole 500mg IV q8h to ceftriaxone or levofloxacin.*

Biliary Source HIGH RISK/SEVERE

FIRST LINE

Piperacillin-tazobactam 4.5gm IV q6hr (or extended infusion)

SECOND LINE

(Type 1 β-Lactam Allergy): Levofloxacin 500mg IV q24hr PLUS Metronidazole 500mg IV q8hr +/- Aztreonam 2gm IV q8hr

NOTE *IF previous colonization or concerns for highly resistant GNRs, consider meropenem 1gm IV q8hr as a substitute for piperacillin-tazobactam or additional GNR coverage to levofloxacin.*

Duration of therapy

Uncomplicated: ≤ 24 hours

Non-operative (uncomplicated) management: 5 days

Complicated by delayed clinical response or inadequate source control: 7-14 days

CONSIDERATIONS

- Due to *E.coli* resistance >10%, empiric quinolone use alone is cautioned in high-risk/severe cases and may warrant double-coverage with aztreonam or an aminoglycoside.
- Ampicillin-sulbactam is not recommended for use because of high rates of resistance among community-acquired *E. coli* and *B. fragilis.*
- Source control as defined per IDSA: single procedure or series of procedures that eliminate infectious foci, control factors that promote ongoing infection, and correct or control anastomatic derangements to restore normal physiologic function.
- Empiric coverage of Enterococcus or Candida is NOT recommended for mild-moderate community-acquired intra-abdominal infections.
- Empiric Enterococcal tx is recommended for health-care associated infections with previous cephalosporin therapy, immunocompromised patients, and those with valvular heart disease or prosthetic intravascular materials.
- Bowel injuries from penetrating, blunt, or iatrogenic trauma repaired w/in 12hr or other intraoperative contamination of the operative field by enteric contents should be treated w/ abx for < 24hrs.
- Use of ursodeoxycholic acid and/or antibiotics for the prevention of biliary stent occlusion or infection is NOT routinely recommended.
- Need for antibiotics in mild, outpatient diverticulitis disease remains controversial.
- Aminoglycosides are not recommended for routine use in adults with community acquired intra-abdominal infection because of the availability of less toxic agents demonstrated to be at least equally effective but may be necessary in high risk/severity patients with Type I PCN or Cephalosporin allergy.

REFERENCES

1. *Joint Surgical Infection Society and Infectious Diseases Society of America Guidelines (CID 2010:50)*
2. *Clinical Practice Guidelines for Antimicrobial Prophylaxis in Surgery (ASHP 2013;70(3))*
3. *Trial of short-course antimicrobial therapy for intraabdominal infection (NEJM 2015;372:1996-2005)*

ORGAN SYSTEM:	SYNDROME:
Urinary Tract	Asymptomatic Bacteriuria

SYMPTOMS AND/OR RISK FACTORS

Asymptomatic bacteriuria (ASB) is defined by isolation of a specific quantity of bacteria in an appropriately collected urine specimen (≥10^5 cfu/mL) from an individual WITHOUT signs or symptoms of infection irrespective of the presence of pyuria.

CULTURE & SUSCEPTIBILITY (C&S) INVESTIGATION

Routine C&S is NOT indicated in asymptomatic patients unless screening in pregnancy or prior to urologic procedure with compromise of the urothelial mucosa.

UPDATE: *Recommendations AGAINST screening for/or treating ASB includes the following:*

- infants & children
- healthy premenopausal, nonpregnant women
- healthy post menopausal women
- older, community-dwelling persons even if functionally impaired
- older persons resident in a LTCF
- diabetic patients
- renal transplant surgery > 1 month prior
- nonrenal solid organ transplant
- low-risk neutropenia (ANC > 100 lasting 7 days or less and clinically stable)
- spinal cord injury
- short-term or long-term indwelling urethral catheter velective nonurologic surgery
- patients undergoing implantation or living with a urologic device

RECOMMENDED TREATMENT AND DURATION

Pregnant women: (select one option)

- Nitrofurantoin 100mg PO BID x 5d
  NOTE: *contraindicated at 38-42 weeks gestation*
- Cephalexin 500mg PO BID x 5d

Urologic procedure:

Direct treatment based on pre-procedure screening C&S. Only 1-2 doses 30-60 minutes before procedure is indicated.

CONSIDERATIONS

- If older patient with functional and/or cognitive impairment with bacteriuria and delirium WITHOUT signs or symptoms of local or systemic infection, assess for other causes and careful observation without antibiotics.
- Bacteriuria identified on preoperative urine screening for non-urologic procedures (cardiac, ortho, vascular) is not an indication for antibiotics and does not decrease surgical site infections or prevent UTIs.

REFERENCES

1. *Nicolle LE et al. Clinical Practice Guideline for the Management of Asymptomatic Bacteriuria: 2019 Update by the IDSA. CID. Ciy1121.*
2. *Updated Beers Criteria.*

ORGAN SYSTEM:	SYNDROME:
Urinary Tract	**Adult Acute Cystitis**

SYMPTOMS AND/OR RISK FACTORS

General symptoms: Acute onset dysuria, frequency or urgency

NOTE: *Consider deviation from the below recommendations (or consult ID) if any of the following risk factors for multidrug resistant organisms are present: antibiotic exposure within 90 days, presence of urinary invasive device(s), history of UTI with multi-drug resistant organism.*

CULTURE & SUSCEPTIBILITY (C&S) INVESTIGATION

If patient requires inpatient admission for acute cystitis, urine C&S are critical in order to optimize therapy.

Urine cultures should be collected from a midstream void prior to antibiotics or a freshly placed urinary catheter.

RECOMMENDED TREATMENT AND DURATION

FIRST LINE: (SELECT ONE OPTION)

- Nitrofurantoin 100mg PO BID x 5d
- Cephalexin 500mg PO BID x 7d

UPDATE: *Fluoroquinolone FDA Safety Alert: Disabling & potentially permanent adverse effects outweigh benefit in cystitis. Only use when no other alternatives exist.*

SECOND LINE:

- Ciprofloxacin 250mg PO BID x 3d

CONSIDERATIONS

- If at risk for STIs w/ symptoms of urethritis, consider screening for chlamydia.
- Scope of this guideline is limited to immunocompetent adults >18 y/o without history of renal transplant.
- For ESBL (Extended Spectrum Beta-lactamase) producing organisms, treat according to reported susceptibility with nitrofurantoin, TMP/SMX or ciprofloxacin. If resistant to all tested antibiotics or multiple allergies, consider Fosfomycin 3gm PO once if available or consult Infectious Diseases for potential alternatives.
- Nitrofurantoin is contraindicated for CrCl < 30mL/min and in pregnancy at term (38-42wks).
- Patients with recurrent UTIs should have empiric therapy selected based upon prior C&S results.
- Chronic antibiotic prophylaxis for most patients with risk factors for recurrent, complicated UTI is NOT typically recommended. Risk of resistance outweighs the slight reduction in infection rate.

NOTE: *One randomized trial confirmed that pre-menopausal women with recurrent UTIs who drank more water (1.5L total fluid daily) got fewer UTIs.*

REFERENCES

1. *Datta R et al. Nitrofurantoin vs fosfomycin. JAMA 2018;319:1771.*
2. *Executive Summary: International Clinical Practice Guidelines for the Treatment of Acute Uncomplicated Cystitis and Pyelonephritis in Women: CID 2011;52(5):561–564.*
3. *2015 Updated Beers Criteria.*
4. *Hooton TM et al. JAMA Intern Med 2018;178(11):1509-1515.*

ORGAN SYSTEM:	SYNDROME:
Urinary Tract	Adult Pyelonephritis

SYMPTOMS AND/OR RISK FACTORS

Upper UTI is frequently associated with general symptoms PLUS back/flank pain, fever & chills.

NOTE: *Consider deviation from the below recommendations (or consult ID) if any of the following risk factors for multidrug resistant organisms are present: antibiotic exposure within 90 days, presence of urinary invasive device(s), history of UTI with multi-drug resistant organism.*

CULTURE & SUSCEPTIBILITY (C&S) INVESTIGATION

If patient requires inpatient admission for acute pyelonephritis, urine C&S are critical in order to optimize therapy.

Urine cultures should be collected from a midstream void prior to antibiotics or a freshly placed urinary catheter.

RECOMMENDED TREATMENT AND DURATION

Inpatient:

FIRST LINE

- Ceftriaxone 1g IV Q24H

SECOND LINE

- Ciprofloxacin 400mg IV Q12H, OR
- Levofloxacin 750mg IV Q24H

Outpatient:

FIRST LINE

- Ceftriaxone 1g IM/IV x 1 dose

  If severe or life-threatening beta-lactam allergy consider Gentamicin 5mg/kg IM/IV x 1 dose

After IM/IV dose of Ceftriaxone or Gentamicin, provide one of the following:

FIRST LINE

- Cephalexin 1g PO TID x 10-14d

SECOND LINE:

- Ciprofloxacin 500mg PO BID x 7d

NOTE: *Above recommendations are for empiric antimicrobial therapy, tailor maintenance therapy to C&S report.*

Duration:

- Duration may vary based upon final antibiotic selection (ex. Cipro 7 days, Levo 5 days, cephalosporin up to 10-14 days)
- GNR bacteremia from a urinary source can safely be treated for 7 days in stable patients without fever.

CONSIDERATIONS

If at risk for STIs w/ symptoms of urethritis, consider screening for chlamydia.

Scope of this guideline is limited to immunocompetent adults >18 y/o without history of renal transplant.

Statewide *E. coli* susceptibility to TMP/SMX is <80% and should be avoided as empiric therapy, but may be considered if confirmed by C&S for pyelonephritis (2 week duration).

Patients with recurrent UTIs should have empiric therapy selected based upon prior C&S results.

Continued ›

For ESBL (Extended Spectrum Beta-lactamase) producing organisms, treat according to reported susceptibility with TMP/SMX or ciprofloxacin. If resistant to all tested antibiotics or multiple allergies, consult Infectious Diseases for potential alternatives. ESBL pyelonephritis may require inpatient admission for IV carbapenem antibiotic.

Chronic antibiotic prophylaxis for most patients with risk factors for recurrent, complicated UTI is NOT typically recommended. Risk of resistance outweighs the slight reduction in infection rate.

UPDATE: *Consider screening patients older than 50 for urogenital cancer after a first episode of pyelonephritis if additional risk factors exist like smoking or obesity.*

REFERENCES

1. *Executive Summary: International Clinical Practice Guidelines for the Treatment of Acute Uncomplicated Cystitis and Pyelonephritis in Women: CID 2011;52(5):561–564.*
2. *Yahav et al. 7 vs. 14 days of antibiotic therapy for uncomplicated gram-negative bacteremia: A non-inferiority randomized controlled trial. CID 2018.*
3. *Kutob LF et al. Effectiveness of oral antibiotics for definitive GNR infections. Intern J Antimicrob Agents 2016;48:498-503.*
4. *Sogaard KK et al. Pyelonephritis persons > 50 clinical marker urogenital cancer. Clin Micro Infect 2018;[e-pub].*

ORGAN SYSTEM:	SYNDROME:
Urinary Tract	**Adult Complicated UTI or Catheter-Associated UTI (CAUTI)**

SYMPTOMS AND/OR RISK FACTORS

Complicated UTI: Infection in males or in the presence of an anatomic/functional abnormality (e.g. enlarged prostate, calculi, obstruction, catheter or stent, neurogenic bladder, neutropenia).

Consider deviation from the below recommendations (or consult ID) if risk factors for multidrug resistant organisms are present.

For long-term care or nursing home residents with altered mental status changes, foul smelling urine, or change in urine color, seek alternative causes (ie. dehydration, medications, environmental changes, metabolic problems, bleeding, stroke). Provide increased fluids (if not contraindicated) and increase monitoring of I/Os and vitals.

CULTURE & SUSCEPTIBILITY (C&S) INVESTIGATION

If patient requires inpatient admission for complicated UTI, urine C&S are critical in order to optimize therapy.

Urine cultures should be collected from a midstream void prior to antibiotics or a freshly placed urinary catheter.

RECOMMENDED TREATMENT AND DURATION

Inpatient:

FIRST LINE:

- Ceftriaxone 1g IV Q24H

SECOND LINE:

- Ciprofloxacin 400mg IV Q12H, OR
- Levofloxacin 750mg IV Q24H

Continued 〉

RECOMMENDED TREATMENT AND DURATION *Continued*

NOTE: *recommendations are for empiric antimicrobial therapy, tailor maintenance therapy to C&S report.*

Outpatient:

Base empiric treatment on prior culture data. If stable vitals & afebrile, provide definitive therapy when new C&S result.

Duration:

- Shorter courses (7 days) are reasonable, if symptoms promptly resolve.
- Longer courses (10-14 days) if delayed response, regardless if catheterized or not.
- If female and < 65 years of age, a 3-day regimen may be considered for CAUTI with catheter removal.

CONSIDERATIONS

Scope of this guideline is limited to immunocompetent adults >18 y/o without history of renal transplant.

Statewide *E. coli* susceptibility to TMP/SMX is <80% and should be avoided as empiric therapy, but may be considered if confirmed by C&S for complicated UTI.

For ESBL producing organisms, treat according to reported susceptibility with TMP/SMX or ciprofloxacin. A carbapenem may be required.

REFERENCES

1. *Diagnosis, Prevention, and Treatment of Catheter-Associated Urinary Tract Infection in Adults: CID 2010; 50:625–663.*

ORGAN SYSTEM:	SYNDROME:
Urinary Tract	**Pediatric FEBRILE Urinary Tract Infection** (Ages 2-24 months)

SYMPTOMS AND/OR RISK FACTORS

Symptoms

Fever, Poor feeding, Vomiting, Irritability, Strong-smelling urine

Diagnostic Criteria for Acute Pyelonephritis

Urinalysis results that suggest infection

- Positive nitrite OR
- Leukocyte esterase OR
- Pyuria AND
- >50,000 CFUs per mL of a uropathogen cultured from a urine specimen obtained through catheterization or SPA

Risk Factors in the absence of another source of infection

Girls: Age <12 months, Temp >39 C, Fever >2 days

Boys: Temp >39 C, Fever >24 hours, Uncircumcised

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Obtain urine culture PRIOR to starting antibiotics

Adjust therapy based on C&S results

RECOMMENDED TREATMENT AND DURATION

Ambulatory Empiric Treatment

FIRST LINE

Cephalexin 50mg/kg/day PO divided TID or QID (max 4gm/day)

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

SECOND LINE (β-LACTAM ALLERGY)

Sulfamethoxazole/trimethoprim 4-5mg/kg PO BID (trimethoprim component for dosing; max 160mg trimethoprim/dose)

Inpatient Empiric Treatment

FIRST LINE

Ceftriaxone 50mg/kg IV Q24H (max 2gm/day)

SECOND LINE (β-LACTAM ALLERGY)

Gentamicin 5mg/kg/day IV

Duration of therapy for either ambulatory or inpatient: 7-10 days

CONSIDERATIONS

- Obtain renal/bladder ultrasound for 1st febrile UTI
- VCUG for 2nd febrile UTI or if abnormalities seen on renal/bladder ultrasound
- If child has received TMP/SMX previously, consider alternative if second line therapy is considered.
- For children > 24 months consider verbal reports of frequency, dysuria, hesitancy, urgency, abdominal/flank pain. Review prior C&S for guidance on empiric treatment if prior history of UTI. It is reasonable to follow same treatment and duration recommendations outlined here.

REFERENCES

(Adopted from the 2018 Alaska Antimicrobial Stewardship Collaborative guide)

1. *Roberts KB. Urinary tract infection: clinical practice guideline for the diagnosis and management of the initial UTI in febrile infants and children 2 to 24 months. Pediatrics. 2011;128(3):595-610.*
2. *Shaw K, et al. Pathway for the Evaluation and Treatment of Children with Febrile UTI. Children's Hospital of Philadelphia. https://www.chop.edu/ clinical-pathway/urinary-tract-infection-uti-febrile-clinical-pathway. Accessed Oct 2018.*

ORGAN SYSTEM:	SYNDROME:
Skin and Soft Tissue	**Uncomplicated Cellulitis in Adults**

SYMPTOMS AND/OR RISK FACTORS

Complicating Risk Factors:

NOTE: *Guideline recommendations are for uncomplicated cellulitis in adults and excludes those with complicating risk factors; if complicating risk factors, treatment may vary and formal ID consultation should be considered.*

- Infected diabetic or vascular ulcer
- Critical illness
- Concern for necrotizing fasciitis
- Deep tissue infection
- Surgical site infection
- Injection drug use
- Human or animal bite
- Bacteremia
- Periorbital or orbital cellulitis
- Perineal/vulvar/perianal infection
- Pregnancy

Diagnostic Studies:

- The following are NOT routinely indicated for initial management of uncomplicated disease: ESR, CRP, Procalcitonin, blood cultures, wound swab/superficial cultures, fungal or AFB cultures, plain films, CT or MRI
- Blood cultures if systemically ill, diabetic or other immunosuppression
- Plain film only if concern for foreign body or necrotizing fasciitis
- Wound culture of purulent drainage

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

- Non-purulent cellulitis is most commonly attributed to beta-hemolytic streptococci
- Purulent cellulitis or cutaneous abscess is most commonly attributed to beta-hemolytic Streptococci or *Staphylococcus aureus* and warrants empiric coverage for MRSA.
- Recurrent MRSA infections need not be cultured at every presentation.
- Gram-negative or anaerobic coverage is usually unnecessary for purulent or non-purulent uncomplicated cellulitis.
- Surgical site infections are dependent upon the site of surgery/ infection.

RECOMMENDED TREATMENT AND DURATION

Non-purulent cellulitis:

FIRST LINE INPATIENT

Cefazolin 2 gm IV q8hr

SECOND LINE INPATIENT

β-Lactam Allergy: Clindamycin 600 mg IV q8hr

FIRST LINE OUTPATIENT or oral step-down

Amoxicillin 500mg PO TID or Cephalexin 500mg-1gm PO TID

SECOND LINE OUTPATIENT

β-Lactam Allergy: Clindamycin 300-450mg PO TID

Purulent cellulitis or cutaneous abscess:

FIRST LINE ADULT INPATIENT

NOTE: *I&D is of utmost importance*

Vancomycin 1gm IV q12hr

FIRST LINE ADULT OUTPATIENT or oral step-down based upon C&S

NOTE: *I&D is of utmost importance*

Antibiotics may not be necessary for drained abscess without surrounding induration or erythema.

TMP/SMX DS 1 tab PO BID or Clindamycin 300mg PO TID or Doxycycline 100mg PO TID

Duration of antibiotics for uncomplicated cellulitis in adults is usually 5 days for uncomplicated cases including a well-drained abscess without surrounding cellulitis but may be extended for severe or poorly responsive disease.

NOTE: *Ibuprofen 600mg PO TID should be added to all situations of cellulitis if no contraindications to NSAID therapy exist.*

CONSIDERATIONS

Antibiotics with broad-spectrum gram-negative activity are NOT recommended except necrotizing fasciitis, and in most cases should be avoided.

Elevate affected area(s).

May consider oral de-escalation options and clinically improving in 2-3 days. Utilize suggested empiric oral options when culture negative or not available.

Treat tinea pedis if applicable.

REFERENCES

1. *Ko LN et al. Imaging & blood cultures in cellulitis. JAMA Intern Med 2018; [e-pub].*
2. *Stevens DL, et al. Practice Guidelines for the Diagnosis and Management of SSTI: 2014 Update by IDSA. CID. 2014; 59(2):e10-e52.*

ORGAN SYSTEM:	SYNDROME:
Skin and Soft Tissue	**Adult Diabetic Foot Infection**

SYMPTOMS AND/OR RISK FACTORS

Assessment

- Physical examination to assess for evidence of infection and depth
- CBC, pre-albumin, BUN, creatinine, hemoglobin A1C, CRP and ESR
- Ankle brachial index (ABI) and/or transcutaneous oxygen tension measurement
- Plain film to assess for foreign bodies, deformity, boney destruction, soft tissue gas, and/or foreign bodies.

NOTE: *metal probe has a negative predictive value of 98% for osteomyelitis; plain film has a specificity 67%, sensitivity 60%*

- When more specific imaging is needed to evaluate for either soft tissue abscess or osteomyelitis an MRI is preferred

Osteomyelitis Evaluation:

- Consider osteomyelitis in any infected, deep, or large foot ulcer, particularly those that are chronic and over bony prominences
- Plain films along with the probe to bone test are reasonable first steps in evaluating for osteomyelitis
- Patients where the diagnosis remains unclear should undergo MRI
- Patients with findings suggestive of osteomyelitis should undergo debridement with bone culture before antibiotics are started if possible
- Consult orthopedics or vascular surgery for potential surgical intervention
- If debridement is not an option an IR guided bone biopsy should be obtained to determine the microbial etiology
- Consult infectious diseases for evaluation and management of long-term antibiotics

Risk

- Infection related to ulceration to the bone, ulcers that have been present for longer than 30 days, recurrent trauma and peripheral arterial disease

Diagnostic Criteria

Obvious purulent drainage AND/OR 2 of the following: Erythema, Pain, Tenderness, Warmth, Induration

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

Infected ulcers initially harbor staphylococcus and streptococcus.

With increasing time, depth and size, wounds are colonized and/or infected with multiple organisms, including Gram negatives and anaerobes

- Do not culture a clinically uninfected lesion
- Do obtain an appropriate specimen for culture from INFECTED wounds and before antibiotics are started, if possible
- Cleanse and debride before collection of tissue
- Tissue collection using sterile scalpel or curettage or biopsy from the base
- Aspirate any purulent secretions using sterile needle & syringe
- Do not obtain a specimen by swabbing the wound or wound drainage

RECOMMENDED TREATMENT AND DURATION

MILD: At least 2 of the following are present at the site of the ulcer/wound: Swelling or induration, erythema, tenderness or pain, warmth, purulent drainage

NOTE: *Use oral agents treating gram-positive cocci only (Beta-hemolytic streptococci and MSSA). Consider addition of MRSA active agent if history of MRSA infection/colonization.*

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

FIRST LINE

Cephalexin 1000mg PO TID OR

Amoxicillin-clavulanate 875/125 mg PO BID

If MRSA concern add: Doxycycline 100 mg PO BID or TMP/SMX DS 1 tab PO BID

SECOND LINE

(Severe PCN Allergy): Clindamycin 300 mg PO TID

Duration for mild infections of soft tissue only is 1-2 weeks.

MODERATE: Local infection with or involvement of deeper structures (abscess, osteomyelitis, septic arthritis) or more extensive erythema (>2 cm spread or associated lymphangitis) without systemic signs of inflammation

NOTE: *May use oral or parenteral agents depending on care location and severity of infection. Treat for pathogens as above plus aerobic gram-negatives. Consider addition of MRSA active agent if history of MRSA infection/colonization.*

Oral Options:

FIRST LINE

Amoxacillin-clavulanate 875/125 mg PO BID

If MRSA concern add: Doxycycline 100 mg PO BID or TMP/SMX DS 1 tab PO BID

SECOND LINE

(Severe PCN allergy): Levofloxacin 750 mg PO daily PLUS Doxycycline 100 mg PO BID

IV Options:

FIRST LINE

Ceftriaxone 2g IV daily PLUS Metronidazole 500mg IV q8h OR

Ampicillin/sulbactam 3gm IV q6h OR

Ertapenem 1gm IV daily

If MRSA concern add: Vancomycin 15 mg/kg IV Q12h

SECOND LINE

(Severe PCN Allergy): Levofloxacin 750 mg IV daily PLUS Clindamycin 900 mg IV q8h

Moderate soft tissue only infections may require 1-3 weeks.

SEVERE: As above with systemic signs of infection (fever, tachycardia, leukocytosis, hypotension, sepsis syndrome, necrotizing infection, etc.) Generally life- or limb-threatening.

NOTE: *Increased frequency of polymicrobial infection. Treat gram-positive cocci including MRSA, aerobic gram-negative rods, and anaerobes. Do not include Pseudomonas coverage unless risk factors (water exposure, previous isolation of Pseudomonas). Consult a surgery team in all severe infections.*

FIRST LINE

Vancomycin 15 mg/kg IV q12h PLUS Ceftriaxone 2g IV daily PLUS Metronidazole 500mg IV q8h (PREFERRED) OR

Vancomycin 15 mg/kg IV q12h PLUS Ertapenem 1g daily OR

Vancomycin 15 mg/kg IV q12h PLUS Piperacillin/tazobactam 4.5g IV q8h (or Extended Infusion)

NOTE: *If water exposure: Treat for Pseduomonas replacing ceftriaxone with cefepime 2gm IV q8hr until cultures return.*

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

SECOND LINE

Severe PCN Allergy: Vancomycin 15 mg/kg IV q12h PLUS Aztreonam 2g IV q8h PLUS Metronidazole 500mg IV q8h

Severe soft tissue infections with initial improvement on IV antibiotics can be switched early to highly bioavailable oral agents (FQ, TMP/SMX, linezolid, metronidazole, etc.) for a combined treatment duration of 2-4 weeks.

Antibiotics can be stopped 2-5 days post resection for bone or joint involvement if complete resection of infected tissue is confirmed post amputation.

If residual soft tissue infection exists after complete bone resection IV and oral antibiotics combined typically lasts 1-3 weeks. If residual infected bone an additional 1-3 weeks is recommended.

Extended durations are likely if no surgery or residual dead bone exists.

REFERENCES

1. *Adopted from the Nebraska Medicine Diabetic Foot Infections Institutional Treatment Guidance. [Accessed March 2019]*
2. *Kwon KT et al. Microbiology and Antimicrobial Therapy for Diabetic Foot Infections. Infection & Chemotherapy 2018;50(1):11-20.*
3. *Lipsky BA et al. 2012 IDSA Guideline for the Diagnosis and Treatment of Diabetic Foot Infections. CID 2012;54(12):132-173.*

ORGAN SYSTEM:	SYNDROME:
Skin and Soft Tissue	**Uncomplicated Cellulitis in Pediatrics**

SYMPTOMS AND/OR RISK FACTORS

NOTE: *Guideline recommendations are for uncomplicated cellulitis in children > 44 weeks and excludes those with complicating risk factors; if complicating risk factors, treatment may vary and formal specialty consultation may be warranted.*

Guideline exclusion criteria:

- Hospital-acquired, surgical site & device-associated infections
- Presumed necrotizing fasciitis
- Orbital/periorbital cellulitis
- Immunodeficiency
- Pressure ulcers
- Solitary dental abscess

Risk factors for MRSA:

- MRSA in the patient
- MRSA in the family
- Recurrent boils, pustules, “spider bites”, that required antibiotics, in patient or family

Specialty Consultation Considerations:

- Orthopedics if deep extremity infection (e.g., tenosynovitis, septic arthritis, osteomyelitis)· Deep puncture wound of hand/fingers/feet
- General surgery if peri-anal abscess (within 1cm of anal verge)· Breast abscess· Perineal abscess· Pilonidal cyst· Large or complex abscess
- ENT if neck abscess
- Dental if facial cellulitis of dental origin

Low Risk Criteria:

Simple abscess · Adequate I&D· Age ≥1 year· No fever· Well-appearing· No significant comorbidities· Follow up assured

Inpatient Admit Criteria (any one of the following):

Systemic illness, not tolerating PO, treatment failure on > 48 hrs of appropriate antibiotics, rapidly progressive lesion, pain control/wound care needed,, inadequate follow-up, all < 2 months of age; consider if < 6 months

Diagnostic Studies:

- The following are NOT routinely indicated for initial management of uncomplicated disease: ESR, CRP, Procalcitonin, blood cultures, wound swab/superficial cultures, fungal or AFB cultures, plain films, CT or MRI
- Perform bedside ultrasound unless clearly fluctuant or draining
- If fluctuant or abscess > 1cm on ultrasound, provide sedation/pain control, I&D and wound culture of purulent drainage
- Obtain a CBC, CRP, and blood cultures in children with signs of systemic toxicity, including ill-appearance, rapidly spreading lesions, persistent fevers, and age < 1 year
- Plain film only if concern for foreign body or necrotizing fasciitis

CULTURE AND SUSCEPTIBILITY (C&S) INVESTIGATION

- Non-purulent cellulitis is most commonly attributed to beta-hemolytic streptococci
- Purulent cellulitis or cutaneous abscess is most commonly attributed to beta-hemolytic Streptococci or Staphylococcus aureus and warrants empiric coverage for MRSA.
- Recurrent MRSA infections need not be cultured at every presentation.
- Gram-negative or anaerobic coverage is usually unnecessary for purulent or non-purulent uncomplicated cellulitis.

RECOMMENDED TREATMENT AND DURATION

Non-purulent cellulitis:

INPATIENT

FIRST LINE INPATIENT

Cefazolin 50 mg/kg IV per day q8hr

SECOND LINE INPATIENT (β-Lactam Allergy)

Clindamycin 25-40 mg/kg per day q6-8hr or Vancomycin if systemic toxicity

OUTPATIENT

FIRST LINE OUTPATIENT OR ORAL STEP-DOWN

Cephalexin 25-50 mg/kg per day divided TID or QID

SECOND LINE OUTPATIENT (β-Lactam Allergy)

Clindamycin 25-30 mg/kg per day TID

Purulent cellulitis or cutaneous abscess:

INPATIENT

FIRST LINE INPATIENT:

Clindamycin 10 mg/kg/dose IV q6-8hr (max does range 600-900mg/dose IV)

SECOND LINE INPATIENT:

Vancomycin 15mg/kg/dose IV q6-8hr (initial max 1gm/dose) if systemically ill, failed outpatient clindamycin, or abscess in an area difficult to drain completely

OUTPATIENT

NOTE: *No systemic antibiotics are needed if adequate I&D and low risk*

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

FIRST LINE OUTPATIENT or oral step-down:

Clindamycin 10 mg/kg /dose PO TID (max single dose range 450-600mg/dose)

SECOND LINE OUTPATIENT:

TMP/SMX 4-6 mg/kg/dose trimethoprim PO BID (max 160mg TMP/dose) or doxycycline if > 8 years 2mg/kg/dose PO BID (max 100mg/dose)

Duration of antibiotics for uncomplicated cellulitis in children is usually 7-10 days. May consider shorter durations (5-7 days) for non-severe infections with quick response to therapy or extended to 14 days for severe disease.

CONSIDERATIONS

- Antibiotics with broad-spectrum gram-negative activity are NOT recommended except necrotizing fasciitis, and in most cases should be avoided.
- Tailor antibiotics if culture results are available; utilize suggested empiric oral options when culture negative or not available.
- May consider oral de-escalation options and clinically improving in 2-3 days.
- If no improvement on adequate antibiotics after 48 hours or significant or rapid progression (ie. more than just 1-2 cm beyond margins) at any time, image (U/S preferred) to rule out abscess formation and consider modification to antibiotic therapy.

NOTE: *The development of a new abscess within an area of previous infection while on antibiotics does not in and of itself constitute treatment failure.*

- Reasonable discharge criteria include: Lesion(s) show signs of improvement, tolerating PO, pain controlled, afebrile > 24 hours, F/U assured within 48 hours
- If fresh or saltwater contact, or other special circumstance, discuss with ID
- If worried about palatability or concerns about administration exist, a single oral antibiotic dose may be given prior to discharge.

REFERENCES

(adopted from Seattle Children's Guide: Simple cellulitis / abscess and UCSF Pediatric Guideline: Skin & Soft Tissue Infections)

1. *1.Kilburn SA et al. Interventions for cellulitis and erysipelas. Cochrane Database of Systematic Reviews 2010 (6). DOI:10.1002/14651858.CD004299.pub2.2*
2. *2.Liu C et al. Clinical practice guidelines by the IDSA for the treatment of MRSA infections in adults and children. CID 2011;52:1-38*
3. *3.Robinson JL et al. Canadian Paediatric Society Infectious Diseases CA-MRSA skin abscesses in children. Paediatr Child Health 2011;16(2):115-64.*
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ORGAN SYSTEM:	SYNDROME:
Miscellaneous	Simplified Approach to Antibiotic Selection in Adults with Severe Sepsis or Septic Shock

ANTIMICROBIAL STEWARDSHIP KEY POINTS

1. Patients presenting with severe sepsis/septic shock are often infected with the same bacteria that cause less severe presentations.
2. The key decision is whether to use an antibiotic (or antibiotic combination) that is based on the specific syndrome (ex. pneumonia or UTI) or to treat sepsis (severe or shock) as an undifferentiated disease state.
3. The most likely pathogens should be covered with the most effective and potent antibiotics. For example, S. pneumoniae is killed very effectively with ceftriaxone.
4. The risk for specific organisms or for drug-resistant infections can be determined by reviewing available data and focusing on the presenting syndrome.

SYMPTOMS AND/OR RISK FACTORS

How much room do you have to be wrong?
Is the patient in acute care or critical care?
Sepsis, severe sepsis or septic shock?

Is the patient at risk of MRSA based upon prior infections, surveillance cultures, risk groups (ie. IVDU, currently incarcerated)?

Should anaerobes be covered based upon extra-biliary colonic source, cavitary aspiration pneumonia?

What risks exist for Pseudomonas (ie. prior *P. aeruginosa* infections, skilled nursing facility or long-term acute care hospital resident)?

RECOMMENDED TREATMENT AND DURATION

If shock, rapid initiation of early broad-spectrum antibiotics as an undifferentiated disease state are warranted.

FIRST LINE ADULT

Meropenem 1gm IV q8hr PLUS Vancomycin

If a syndrome based approach to sepsis or severe sepsis, consider the following key agents for adequate empiric coverage based upon risk of MRSA, anaerobes or pseudomonas.

If Risk of MRSA

FIRST LINE ADULT

Include Vancomycin IV loading dose X 1 (2gm if ≥ 70kg, 1.5gm if < 70kg) STAT, then 15mg/kg IV q12hrs

If Risk of anaerobes

FIRST LINE ADULT

Include Metronidazole 500mg IV q8hr

If Risk of pseudomonas:

FIRST LINE ADULT

Include Cefepime 2gm IV q8hr

If Risk for highly resistant gram-negative pathogens including Acinetobacter or ESBLs:

FIRST LINE ADULT

Include Ciprofloxacin 400mg IV q8hr

SECOND LINE ADULT

Include Tobramycin 7mg/kg IV q24hr

NOTE *Antibiotic recommendation assumes that these drugs or spectrum of activity are not already included in the syndrome-based approach to sepsis.*

Continued ›

RECOMMENDED TREATMENT AND DURATION *Continued*

CNS CONSIDERATIONS

If adult patient presents with concerns for meningitis, ensure adequate coverage for *S. pneumoniae, N. meningitidis* and *H. influenzae*; consider Listeria and HSV in patients age > 50, immunocompromised or alcoholic)

Order antibiotics immediately.

LP for opening pressure, gram stain, culture, HSV PCR, cell count, glucose and protein.

Do not wait for results of LP to initiate antimicrobials.

Add cryptococcal antigen for HIV patients.

Non-surgical community-acquired meningitis:

FIRST LINE:

Ceftriaxone 2gm IV q12hr PLUS Vancomycin IV loading dose X 1 (2gm if ≥ 70kg, 1.5gm if < 70kg) STAT, then 15mg/kg IV q8hrs

ADD: Ampicillin 2gm IV q4hr for Listeria coverage

ADD: Acyclovir 10mg/kg IV q8hr for HSV coverage

Consider dexamethasone 0.15mg/kg IV q6hr for 2-4 days, give 15 minutes prior to antibiotics if possible

Post-surgical meningitis:

S. epidermidis, S. aureus, P. acnes, gram-negative rods (including P. aeruginosa) should be covered empirically.

FIRST LINE:

Cefepime 2gm IV q8hr PLUS Metronidazole 500mg IV q8hr PLUS Vancomycin IV loading dose X 1 (2gm if > 70kg, 1.5gm if < 70kg) STAT, then 15mg/kg IV q8hrs

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