



PeaceHealth Empirical Antimicrobial Treatment Guide

| Site of infection | | Likely pathogens | Empirical treatment of choice | Standard duration of therapy | Comments |
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| Urinary tract | Asymptomatic bacteriuria | Any bacteria, regardless of colony count or presence of pyuria, LE, nitrite, etc. | Antimicrobials should not routinely be prescribed for asymptomatic bacteriuria | 0 days | Patients who are pregnant, with invasive GU surgery, or febrile neutropenia may require antibiotics |
| | Asymptomatic bacteriuria and altered mental status | | Do not treat asymptomatic patients with delirium/dementia unless sepsis with fever or leukocytosis, with no other source identified, and where UTI is not ruled out via absence of pyuria. | See duration for pyelonephritis (cystitis excluded if UTI with sepsis; fever +/- leukocytosis) | Consider imaging to aid diagnosis of UTI with sepsis; fever/leukocytosis absent GU symptoms in patients with delirium or dementia |
| | Cystitis | <i>E. coli</i> , <i>Klebsiella</i> spp., <i>Proteus</i> spp., other enterobacterales. | Nitrofurantoin 100 mg PO Q12H -OR- SMX/TMP DS PO Q12H -OR- Cephalexin 1-3 g/day PO in 2-3 doses -OR- Tobramycin 5 mg/kg x1 | 5 days 3 days 5 days 1 dose | Nitrofurantoin only for uncomplicated cystitis and estimated CrCl \geq 40 mL/min Gentamicin 5 mg/kg x1 adequate if not <i>P. aeruginosa</i> |
| | Pyelonephritis or UTI with signs of systemic illness | | Ceftriaxone 1 g IV Q24H If suspicion for resistant pathogen: Cefepime 1g IV Q8H -OR- Piperacillin/tazobactam 3.375 IV Q8H | 7 days (10 total if SMX/TMP or oral β -lactam) | Broader spectrum should only be used for patients with hemodynamic instability and/or prior relevant cultures demonstrating resistance |
| Lungs | Community acquired pneumonia (CAP) | <i>S. pneumoniae</i> , <i>H. influenzae</i> , <i>S. aureus</i> , <i>M. catarrhalis</i> ; rarely atypical organisms | Ceftriaxone 1 g IV Q24H +/- Azithromycin 500 mg PO Q24H | 5 days | Addition of coverage for MRSA or <i>P. aeruginosa</i> should be avoided without a history of prior relevant cultures |
| | Aspiration pneumonia | Oral anaerobes | Ceftriaxone 1 g IV Q24H -OR- Ampicillin/sulbactam 3g IV Q6H | 5 days | Aspiration events or aspiration pneumonitis should not be treated empirically. For aspiration pneumonia, metronidazole is not needed |
| | Hospital acquired pneumonia (HAP) | Above plus MRSA, enterobacterales, <i>P. aeruginosa</i> | Cefepime 2g IV Q8H +/- Vancomycin per pharmacy | 7 days | Pending cultures, de-escalate as soon as possible. MRSA nares PCR has excellent NPV |

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For complex infections please consider infectious diseases consultation

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| Skin | Cellulitis (non-suppurative) | <i>S. pyogenes</i> (Group A strep), <i>S. agalactiae</i> (Group B strep) | Cefazolin 2 g IV Q8H | 5 days | Streptococcal cellulitis may appear to worsen for 24-48 hours on appropriate therapy; this is an expected finding rather than an indication to broaden therapy |
| | Abscess | <i>S. aureus, viridans Streptococcus</i> spp. | Incision and drainage; consider: Cefazolin 2 g IV Q8H -OR- Vancomycin per pharmacy | 0-5 days Based on pathogen and clinical status | I&D alone may be sufficient; short course oral antibiotics indicated for drained abscess with surrounding cellulitis |
| | Necrotizing skin and soft tissue infections | <i>S. pyogenes</i> (Grp. A strep), <i>S. aureus</i> , <i>E. coli</i> , anaerobes | Piperacillin/tazobactam 3.375 g IV Q8H OR Ceftriaxone 1 g IV Q24H + metronidazole 500 mg IV Q12H if penicillin allergic + vancomycin per pharmacy +/- clindamycin 600 – 900 mg IV Q8H or linezolid 600 mg IV Q12H | Improvement; afebrile for 48-72 hours | If hemodynamically stable, clindamycin not needed. If linezolid for streptococcal toxin suppression, vancomycin not needed. De-escalate based on cultures |
| | Diabetic foot infections (including those with osteomyelitis) | <i>Staphylococcus</i> spp., <i>Streptococcus</i> spp., enterobacteriales, anaerobes | Cefazolin 2g IV Q8H OR Ampicillin/sulbactam 3 g IV Q6H OR Ceftriaxone 1 g IV Q24H If ischemic limb, necrosis, gas forming: +/- Metronidazole 500 mg IV or PO Q12H If increased risk of <i>P. aeruginosa</i> * Cefepime 1-2 g IV Q8H OR Piperacillin/tazobactam 3.375 g IV Q8H If increased risk for MRSA* add Vancomycin per pharmacy | 7-14 days If adequate surgical debridement: 10 days If osteomyelitis with surgical source control and positive margin culture: 14-21 days | * <i>P. aeruginosa</i> is an unusual pathogen in diabetic foot infections. Increased risk of clinically relevant <i>P. aeruginosa</i> with positive culture within 3 weeks or macerated wound. De-escalate as soon as possible. *MRSA risk increased with history of MRSA wound infection Ischemic limb, necrosis, or gas are NOT indications for anti-pseudomonal or MRSA activity |
| | Bite wounds | <i>Staphylococcus</i> spp., <i>Streptococcus</i> spp., oral anaerobes including <i>Pasteurella multocida</i> , HACEK organisms | Ampicillin/sulbactam 3 g IV Q6H OR Ceftriaxone 1 g IV Q24H +/- Metronidazole 500 mg IV or PO Q12H | 7 days | HACEK: <i>Haemophilus</i> , <i>Aggregatibacter</i> , <i>Cardiobacterium</i> , <i>Eikenella</i> , <i>Kingella</i> spp. |

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| Abdomen | Variable | <i>Viridans streptococcus</i> spp., enterobacterales, anaerobes including <i>B. fragilis</i> | Ceftriaxone 1 g IV Q24H + metronidazole 500 mg IV/PO Q12H If suspicion for resistant pathogens*: Cefepime 1-2g IV Q8H + metronidazole 500 mg IV/PO Q12H -OR- Piperacillin/tazobactam 3.375 g IV Q8H | 4 days from surgical source control, otherwise dependent on clinical status | No indication for prophylactic antimicrobials for pancreatitis, even with necrosis, unless confirmed infection present *Biliary infections with anastomosis *Post-operative infections |
| GI tract | Odontogenic/oral | <i>Staphylococcus</i> spp., <i>Streptococcus</i> spp., oral anaerobes including <i>Pasteurella multocida</i> , HACEK organisms | Ampicillin/sulbactam 3g IV Q6H Ceftriaxone 1g IV Q24H +/- metronidazole 500 mg IV/PO Q12H Amoxicillin/clavulanate 875 mg PO BID | Dependent on source control, typically no more than 7 days | <i>Pseudomonas</i> should not be treated for empirically in odontogenic infections. |
| | Infectious colitis with bloody diarrhea | <i>Campylobacter</i> , shiga-toxin producing <i>E. coli</i> (STEC), <i>Salmonella</i> , <i>Shigella</i> spp. | Avoid empiric antibiotics without signs of severe sepsis due to added risk of hemolytic uremic syndrome. Ceftriaxone 1g IV Q24H (<i>Salmonella</i> or <i>Shigella</i> spp.) -OR- Azithromycin 500 mg IV/PO Q24H (<i>Campylobacter</i> or <i>Shigella</i> spp.) | 3-7 days depending on pathogen/site | Antibiotics do not alter and may worsen illness in many cases. Consider antibiotics with pathogen identification for non-STEC in immune compromised or those with severe disease. Consider ID consult for bloodstream infections with <i>Salmonella</i> spp. |
| | <i>Clostridioides difficile</i> (formerly <i>Clostridium difficile</i>) | <i>Clostridioides difficile</i> (formerly <i>Clostridium difficile</i>) | Vancomycin 125 mg PO Q6H Fulminant: vancomycin 500 mg Q6H, oral AND/OR rectal +/- metronidazole 500 mg IV Q8H | 10 days | Fulminant: hypotension or shock, ileus, megacolon attributable to <i>C. difficile</i> (rare) |
| | <i>H. pylori</i> | <i>H. pylori</i> | Tetracycline 500 mg PO Q6H Metronidazole 500 mg PO Q6-8H Pantoprazole 40 mg PO Q12H Bismuth subsalicylate 262 mg PO Q6H | 14 days | Other agents not recommended without susceptibility data |
| | SBP (confirmed by PMN or empirical in the setting of GI bleed and ascites) | <i>Enterobacterales</i> , <i>Viridans streptococcus</i> | Ceftriaxone 1g IV q24H | 5 days (varices) 5-7 days (confirmed) | Stopping at resolution of hemorrhage, hospital discharge, or for PMN less than 250 on repeat sampling if less than 5 days is non-inferior for variceal bleeding |
| CNS | Meningitis | <i>S. pneumoniae</i> , <i>N. meningitidis</i> , <i>L. monocytogenes</i> | Ceftriaxone 2 g IV Q12H + Vancomycin per pharmacy | <i>N. meningitidis</i> : 7 days <i>S. pneumoniae</i> : | Ampicillin indicated for adults age > 50 or patients who are immune compromised or |

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| | | | +/- Ampicillin 2 g IV Q4H Dexamethasone 10 mg IV Q6H | 14 days <i>L. monocytogenes</i> : 21 days | pregnant. If started with or before antibiotics, continue dexamethasone for <i>S. pneumoniae</i> , consider benefit v. risk for <i>H. influenzae</i> , and stop if <i>L. monocytogenes</i> |
| | Encephalitis | HSV-1 and -2, VZV | Acyclovir 10 mg/kg IV Q8H | 14-21 days | No specific treatment is recommended for viral meningitis |
| Musculoskeletal | Discitis or osteomyelitis | <i>S. aureus</i> , coagulase negative <i>Staphylococcus</i> spp., enterobacterales | Hold empiric antibiotics absent clinical suspicion for bacteremia; if indicated: Vancomycin per pharmacy +/- Ceftriaxone 2g IV Q24H | 4-6 weeks | Empiric antibiotics decrease diagnostic yield of cultures if bacteremia not present. Please consult ID |
| | Septic arthritis | <i>S. aureus</i> , <i>Streptococcus</i> spp., <i>N. gonorrhoeae</i> | Hold empiric antibiotics absent clinical suspicion for bacteremia; if indicated: Vancomycin per pharmacy +/- Ceftriaxone 2g IV Q24H See recommendations by likely source, with hemodynamic instability if present | 2-4 weeks | Empiric antibiotics decrease diagnostic yield of cultures if bacteremia not present. Please consult ID Pending cultures, de-escalate as soon as possible |
| Severe sepsis | See infections by likely source | See infections by likely source | See recommendations by likely source, with hemodynamic instability if present | See infections by likely source | Pending cultures, de-escalate as soon as possible |
| Cardiovascular | Infective endocarditis | <i>Streptococcus</i> spp., <i>S. aureus</i> (MRSA with relevant clinical history (IVDU), HACEK orgs. | If unstable: Ceftriaxone 2 g IV Q24H +/- Vancomycin per pharmacy | 2-6 weeks | Empiric therapy not needed if stable; definitive treatment based on cultures appropriate, please consult ID |