



The use of the following guidelines for empiric therapy should include patient-specific information, including drug allergies and renal/liver function; decisions should be based on the judgment of the clinician. Definitive therapy should be used when culture results/data are available.					
Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment		Ref.
<b>Central Nervous System</b>					
<a href="#">Bacterial meningitis</a>	Neonatal	<i>S. agalactiae</i> (GBS) <i>L. monocytogenes</i> <i>E. coli</i> <i>Klebsiella spp.</i>	Ampicillin + Ceftazidime if <4 weeks corrected age  * Pediatric ID consult recommended (Consideration in the ddx of neonatal viral infections and/or TORCH infections)	<i>S. agalactiae</i> (GBS) 14-21 days <i>L. monocytogenes</i> ≥ 21 days Aerobic gram-negative bacilli (e.g., <i>E. coli</i> , <i>Klebsiella spp.</i> ) <sup>†</sup> 21 days	(1), (2)
	Ages > 1 month	<i>S. pneumoniae</i> <i>N. meningitidis</i> <i>H. influenzae</i>	Ceftriaxone + Vancomycin	<i>S. pneumoniae</i> 10-14 days <i>N. meningitidis</i> 7 days <i>H. influenzae</i> 7 days	
Brain abscess	<i>Streptococcus spp.</i> (e.g., <i>S. anginosus</i> group, <i>S. pneumoniae</i> , <i>S. pyogenes</i> ) <i>S. aureus</i> Enterobacterales spp. <i>H. influenzae</i> <i>Anaerobes</i> (e.g., <i>Bacteroides</i> , <i>Fusobacterium spp.</i> ; <i>Prevotella</i> ) <i>Citrobacter spp.</i> (typically neonates)	Ceftriaxone + Metronidazole + Vancomycin  * Pediatric ID, ENT, and Neurosurgical consults recommended	4 – 6 weeks		(3)
<a href="#">Healthcare-associated meningitis/ventriculitis</a>	<i>S. aureus</i> Coagulase-negative staphylococci (CoNS) Aerobic gram-negative bacilli (including <i>P. aeruginosa</i> , multidrug resistant gram-negative organisms) <i>C. acnes</i> (formerly <i>Propionobacterium acnes</i> )	Cefepime + Vancomycin  * Pediatric ID consult recommended	<i>S. aureus</i> or gram-negative bacilli ≥ 10-14 days <sup>†</sup> CoNS or <i>C. acnes</i> (minimal CSF pleocytosis, normal CSF glucose) ≥ 10 days <sup>†</sup> CoNS or <i>C. acnes</i> (significant CSF pleocytosis, low CSF glucose) ≥ 10-14 days <sup>†</sup>	(4)	
			<sup>†</sup> Days after last positive culture Note: Shunt removal is recommended		

Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.	
<b>Ears, Nose, Throat &amp; Respiratory Tract</b>					
<a href="#">Acute otitis media (AOM)</a>	<i>S. pneumoniae</i> <i>H. influenzae</i> <i>M. catarrhalis</i>	Observation (no antibiotics): Use of symptomatic relief with observation and close follow-up is an option for unilateral AOM in children ≥ 6 months of age with non-severe signs or symptoms (i.e., mild otalgia for less than 48 hours and < 39°C [102.2°F], no otorrhea); initiate antibiotic therapy if the child worsens or fails to improve within 48 to 72 hours of symptoms onset.  1 <sup>st</sup> line when antibiotic therapy indicated: Amoxicillin (high dose 80-90 mg/kg/day divided BID, max 4 gm/day)  Receipt of Amoxicillin within last 30 days, concurrent conjunctivitis, or persistent symptoms following 48-72hr of 1 <sup>st</sup> line therapy: Amoxicillin-clavulanate (see <a href="#">link</a> )  If penicillin allergy, Cefdinir 14 mg/kg/day divided BID	< 2 years or severe symptoms*	10 days	(5)
			2 - 5 years w/ mild-moderate symptoms	7 days	
			≥ 6 years w/ mild-moderate symptoms	5-7 days	
			* Severe symptoms in all ages include moderate or severe otalgia or otalgia for at least 48 hours or temperature 39°C [102.2°F] or higher.		
<a href="#">Cervical lymphadenitis</a>	Group A -beta hemolytic streptococci <i>S. aureus</i> (MSSA) Oral anaerobes	Amoxicillin-clavulanate (see <a href="#">link</a> )  If Penicillin allergy, Clindamycin	Uncomplicated or adequate source control: 5-10 days (Tailor duration based on resolution of signs and symptoms.)		(6), (7)
	<i>S. aureus</i> (MRSA)	Trimethoprim-sulfamethoxazole (oral option) OR Vancomycin (IV only) OR Clindamycin, if susceptible			(8)
	Non-tuberculous mycobacteria (e.g., MAC)	Surgical excision typically curative. Start antimycobacterial therapy only if incomplete or no surgical excision or if recurrent disease. Medical treatment controversial, includes macrolide + (Ethambutol AND/OR Rifabutin OR Rifampin)	If antimycobacterial drugs indicated, duration may require >12 weeks		(6), (9)
	<i>Mycobacterium tuberculosis</i>	4 drugs: RIPE – Rifampin + Isoniazid + Pyrazinamide + Ethambutol *Pediatric ID consult recommended	≥ 6 months		(6), (10)
	<i>Bartonella henselae</i> (uncomplicated cat scratch disease)	Azithromycin *Pediatric ID consult recommended for complicated disease (e.g., osteitis, hepatosplenic disease)	5 days		(6)
<b>Epiglottitis</b>	<i>H. influenzae</i> Beta hemolytic streptococci <i>S. pneumoniae</i> <i>S. aureus</i>	Ceftriaxone  If septic appearing, consider adding vancomycin	7 - 10 days		(11), (12)

Diagnosis		Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.
Malignant otitis externa		<i>Pseudomonas spp.</i> <i>S. aureus</i> (less frequent)	Cefepime  If penicillin allergy or step-down oral therapy, ciprofloxacin *Pediatric ID consult recommended	≥ 6 weeks (surgical intervention often indicated)	(13)
		<i>Aspergillus spp.</i>	Voriconazole *Pediatric ID consult recommended	>12 weeks	(14)
Mastoiditis	Acute	<i>S. pneumoniae</i> Beta hemolytic streptococci <i>S. aureus</i> <i>H. influenzae</i> <i>Fusobacterium spp.</i>	Ampicillin-sulbactam  Consider Vancomycin if MRSA risk factors Consider transition to oral regimen (e.g., Amoxicillin-clavulanate, see <a href="#">link</a> ) when clinically appropriate Consider surgical drainage *Pediatric ID consult recommended	Uncomplicated: 10 days  Complicated (including extra- and intracranial complications including subperiosteal abscess or osteomyelitis): 4 weeks	(15), (16),
	Chronic or recurrent AOM	<i>P. aeruginosa</i> Enteric gram-negative bacteria <i>S. aureus</i> <i>S. pneumoniae</i> Anaerobes	Cefepime + Metronidazole + Vancomycin *Pediatric ID consult recommended	4 weeks	(16), (17)
Odontogenic infection	Mild, no abscess	Polymicrobial; Viridans group streptococci <i>Peptostreptococcus</i> <i>Actinomyces</i> <i>Lactobacillus spp.</i>	Amoxicillin 45 mg/kg/day divided TID, max 1500 mg/day  If NPO, Ampicillin If severe beta-lactam allergy, Clindamycin	7 days or until 3 days after symptom resolution (whichever is longer) with surgical intervention	(18 - 20)
	Moderate/severe, abscess	As above and follows: <i>Prevotella spp.</i> <i>Neisseria spp.</i> <i>Fusobacterium spp.</i> <i>Clostridium spp.</i> <i>Staphylococcus spp.</i> <i>Eikenella spp.</i> <b>Caution:</b> Culture results may have poor recovery of all pathogens involved	Ampicillin-sulbactam  If Penicillin allergy (non-severe), Cefazolin + Metronidazole If severe beta-lactam allergy, Ciprofloxacin + Metronidazole	7 - 10 days or until 3 days after symptom resolution (whichever is longer) with surgical intervention *Consider transition IV to oral in patients without bacteremia if surgically controlled and clinical improving	(47 - 51)
	Severe, abscess, with no clinical improvement after trialing therapy listed above	<i>Clostridium spp.</i> <i>Staphylococcus spp.</i> <i>Eikenella spp.</i> <b>Caution:</b> Culture results may have poor recovery of all pathogens involved	Cefepime + Metronidazole + Vancomycin  If severe beta-lactam allergy, Aztreonam + Metronidazole + Vancomycin  *Pediatric ID consult recommended	≥ 10 days or until 3 days after symptom resolution (whichever is longer) with surgical intervention	
Peritonsillar, parapharyngeal, or retropharyngeal abscess		Beta hemolytic streptococci <i>S. aureus</i> (MSSA or MRSA) Respiratory anaerobes Kawasaki disease	Ampicillin-sulbactam  If penicillin allergy, Ceftriaxone + Clindamycin  Add Vancomycin for toxic appearing patients pending culture results	Parapharyngeal or retropharyngeal abscess: 14 days  Submandibular space infections: 2-3 weeks	(21)

Diagnosis		Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment		Ref.
Pertussis		<i>B. pertussis</i> <i>B. parapertussis</i>	Azithromycin  If Azithromycin contraindicated: Trimethoprim-sulfamethoxazole (TMP/SMX) in children ≥ 2 months	Azithromycin x 5 days TMP/SMX x 14 days (avoid in children <2 months)		(22)
Pharyngitis		Beta hemolytic streptococci (Group A streptococci)	Penicillin VK (patients ≤27 kg: 250 mg 2-3 times/day and patients >27 kg: 500 mg 2-3 times/day OR Amoxicillin 50 mg/kg/day once daily, max 1 gm/day  Allergic: Cephalexin or Clindamycin or Azithromycin	10 days  [For Azithromycin PO: <u>12 mg/kg/day</u> (max: 500 mg/dose) X 5 days or 20 mg/kg/day (max: 1000 mg/dose) x 3 days]		(23), (54)
		<i>Arcanobacterium</i>	Azithromycin			
Pneumonia	Aspiration pneumonia	<i>Enteric Gram negative Oral flora</i> <i>Anaerobic bacteria</i>	Ampicillin  If not fully immunized or failed Amoxicillin or Ampicillin, Ceftriaxone  If empyema or lung abscess present, Ampicillin-sulbactam or Ceftriaxone + (Clindamycin or Metronidazole)  If hospital-acquired aspiration pneumonia, Piperacillin-tazobactam  Review previous microbiologic history to guide antibiotic selection	7 - 10 days		(24), (25)
	<a href="#">Community-acquired pneumonia</a> (Ages >3 months)	<i>S. pneumoniae</i> Group A streptococci <i>S. aureus</i> Viruses ( <i>M. pneumoniae</i> , <i>C. pneumoniae</i> if school-aged children and adolescents)	PO: Amoxicillin 90 – 100 mg/kg/day divided BID or TID, max 4 gm/day  If not fully immunized, Amoxicillin-clavulanate (see <a href="#">link</a> )  If beta-lactam allergy, see “Clinical Pathways” for alternative enteral agents  IV: Ampicillin  Alternative: Ceftriaxone if not fully immunized, abscess or empyema present, or critically ill  *For patients > 5 years of age, consider addition of Azithromycin [10 mg/kg (max, 500 mg) PO on day 1, then 5 mg/kg (max, 250 mg) daily on days 2-5] for suspected atypical pneumonia	Uncomplicated 5 days  Complicated (parapneumonic effusion or empyema) Prolonged, typically 2-4 weeks	*Consider transition IV to oral in patients without bacteremia if clinically improved (as early as 2–3 days after the start of IV therapy)	(26), ( )
	<a href="#">Hospital-acquired pneumonia</a>	<i>S. aureus</i> (MRSA) <i>P. aeruginosa</i> Gram-negative bacilli ( <i>E. coli</i> , <i>Klebsiella</i> , <i>Serratia</i> , <i>Enterobacter</i> )	[Cefepime OR Piperacillin-tazobactam OR Meropenem] + Vancomycin (guided by results of MRSA nasal swab)  *Pediatric ID consult recommended	Duration should be guided by improvement in clinical and laboratory parameters, although evidence suggests that <u>7 days</u> of therapy is sufficient in most cases		(27)

Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.
<a href="#">Sinusitis</a>	<i>S. pneumoniae</i> <i>H. influenzae</i> <i>M. catarrhalis</i> <i>S. aureus</i> (dental origin) Anaerobes (chronic)	Uncomplicated, mild infection (age 2 years or older): Amoxicillin (45 mg/kg/day divided BID)  <u>Complicated infection [severe onset and/or worsening course of infection OR age &lt;2 years OR unimmunized OR daycare attendance OR receipt of antibiotics within last 30 days]:</u> Amoxicillin-clavulanate (see <a href="#">link</a> )  If severe penicillin allergy, Cefdinir If severe cephalosporin allergy, Levofloxacin  Severe infection requiring hospitalization: Ampicillin-sulbactam	Uncomplicated, mild infection: 7 days  Complicated sinusitis or requiring hospitalization: 10 - 14 days or 7 days after patient's symptoms resolve (minimum 10 days)	(28), (29), (65), (66)
<b>Eye</b>				
<b>Orbital cellulitis</b>	<i>S. pneumoniae</i> Streptococci <i>S. aureus</i> <i>H. influenzae</i> (unimmunized) <i>M. catarrhalis</i> Anaerobes	Ampicillin/sulbactam  If concern for extension into CNS: Ceftriaxone + Metronidazole + Vancomycin  *Pediatric ID consult recommended, if complicated	Uncomplicated: 2-3 weeks Complicated: ID consult  Transition IV to oral amoxicillin-clavulanate when clinically improved	(30)
<b>Preseptal (periorbital) cellulitis</b>	<i>S. pneumoniae</i> <i>S. aureus</i> Beta hemolytic streptococci <i>H. influenzae</i>	PO: Amoxicillin-clavulanate (see <a href="#">link</a> )  IV: Ampicillin-sulbactam  *If extensive cellulitis or ill appearing patient, consider adding Vancomycin pending culture results	7 - 10 days  Transition IV to oral Amoxicillin-clavulanate when clinically improving	(31)
<b>Heart</b>				
<a href="#">Endocarditis (bacterial)</a>	*Pediatric ID consult recommended			(32)
<b>Blood</b>				
<b>Sepsis</b>	Refer to LPCH "Severe Sepsis/Septic Shock Pathway" per the Clinical Pathways and Target Based Care Website			
<b>Staphylococcal bacteremia</b>  *Pediatric ID consult recommended for <i>S. aureus</i> bacteremia	<i>S. aureus</i> (MSSA or MRSA) <i>S. lugdunensis</i>	Methicillin-sensitive: Cefazolin OR Nafcillin (central line administration preferred)  Methicillin-resistant: Vancomycin  <i>S. aureus</i> has a propensity to form biofilms; line removal is strongly recommended. For patients unable to undergo line removal, lock therapy may be considered on a case-by-case basis in conjunction with ID consult.	Uncomplicated <sup>†</sup>   14 days from first negative culture Complicated   4-6 weeks from first negative culture	(8), (33) (34)
	Coagulase Negative Staphylococcus ( <i>S. epidermidis</i> , <i>S. hominis</i> )	Methicillin-sensitive: Cefazolin OR Nafcillin (central line administration preferred)  Methicillin-resistant: Vancomycin  *Longer durations of antibiotic therapy may be required for indwelling hardware or surgical material present. ID consult recommended	<i>Single set with only one positive bottle may be representative of contamination and not require treatment</i> No catheter present: 7 days Catheter removal: 7 days post-removal Catheter retained: 10 days from 1st negative culture	(33)

Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.
<p><b>GI/GU</b></p> <p><u><a href="#">C. difficile infection</a></u></p> <p>- CDI <b>episode</b>: More than 3 watery bowel movements in 24 hours and a positive stool test for <i>C. difficile</i> toxin by PCR</p> <p>- CDI <b>recurrence</b>: An episode of CDI occurring within 8 weeks of a previous episode</p> <p><i>In general, only toxin-positive (EIA+) patients benefit from C. difficile treatment.</i></p>	<p><b>Initial CDI episode and subsequent episodes &gt; 8 weeks apart</b></p> <p><b>First CDI recurrence</b></p> <p><b>Second or subsequent CDI recurrence</b></p> <p><b>Severe/fulminant CDI (hypotension/shock, ileus, megacolon)</b></p> <p><b><u>Recommend ID consult</u></b></p>	<p>Vancomycin <b>PO</b> <sup>a,b</sup></p> <p><i>Alternative for NPO: Metronidazole IV</i></p> <p>Vancomycin <b>PO</b> <sup>a,b</sup></p> <p>Vancomycin <b>PO</b> + taper <sup>a,b</sup> OR Fidaxomicin (NF) <b>PO</b> x 10 days<sup>a-c</sup></p> <p><i>If fidaxomicin PO 10-day course was used for previous episode: Fidaxomicin (NF) <b>PO</b> tapered course <sup>a-c</sup> --- Taper should start no later than day 5 of therapy.</i></p> <p><b>For 2+ recurrences:</b> Consider Gastroenterology consult for evaluation of Fecal microbiota transplant (FMT)</p> <p>Vancomycin <b>PO</b> <sup>a,b</sup> + Metronidazole <b>IV</b> or <b>PO</b></p> <p>If ileus, consider adding rectal instillation of vancomycin four times daily to regimen. Not appropriate for neutropenic patients. Vancomycin rectal enema: 1-3 years: 250mg (50 mL); 4-9 years: 375mg (75 mL); &gt;10 years: 500mg (100 mL)</p> <p><sup>a</sup> <b>Dosing: General dosing recommendations are available in the HSM.</b> <b>Vancomycin PO taper</b></p> <ul style="list-style-type: none"> <li>• Vancomycin 10 mg/kg (max 125 mg/dose) QID x 10 days, then BID x 7 days, then once daily x 7 days, then daily every 2-3 days x 2-8 weeks.</li> </ul> <p><b>Fidaxomicin (NF) PO:</b></p> <ul style="list-style-type: none"> <li>• Standard course: 16 mg/kg BID (max 200 mg/dose) BID x 10 days</li> <li>• Tapered course: 16 mg/kg BID (max 200 mg/dose) BID x 5 days, then 16 mg/kg (max 200 mg/dose) every other day for 20 days.</li> </ul> <p><sup>b</sup> Enteral administration of vancomycin or fidaxomicin includes PO and gastric tube administration with either the oral suspension or solid dosage formulations (opening vancomycin capsules or crushing fidaxomicin tablets).</p> <p><sup>c</sup> <b>Fidaxomicin is non-formulary and requires Pediatric ID approval prior to inpatient use.</b></p>	<p>10 days</p> <p>If symptoms have improved but not resolved by 10 days, consider extending treatment duration to 14 days.</p> <p>Discontinue any potentially offending agents, including antibiotics, as soon as is clinically appropriate. In patients requiring systemic antibiotics during or shortly after CDI treatment, some experts recommend continuing oral vancomycin until completion of systemic antibiotics.</p>	<p>(35, 58-64)</p>

Diagnosis	Usual Etiology	Suggested Empiric Therapy		Usual Duration of Treatment		Ref.		
<b>Intra-abdominal infection (including cholecystitis, necrotizing enterocolitis and perforated appendicitis)</b>	<i>E. coli</i> <i>Klebsiella spp.</i> <i>Pseudomonas</i> <i>Proteus</i> <i>Enterobacter spp.</i> Anaerobes (e.g., <i>B. fragilis</i> ) Streptococci <i>Enterococcus spp.</i>	Uncomplicated	Ceftriaxone + Metronidazole	Uncomplicated/ Complicated	4-7 days	(36)		
		Complicated/ healthcare-associated	([Cefepime or Ceftazidime or Ciprofloxacin] + Metronidazole ± Ampicillin*) <u>OR</u> Piperacillin-tazobactam <u>OR</u> Meropenem				Cholecystitis	4-7 days
		Necrotizing enterocolitis (NEC) - use vancomycin instead of ampicillin for suspected MRSA or ampicillin-resistant enterococcus	([Ampicillin or Vancomycin] + [Gentamicin or Ceftazidime] AND Metronidazole if perforation) <u>OR</u> Piperacillin-tazobactam <u>OR</u> Meropenem				Necrotizing enterocolitis	7-10 days
		Perforated appendicitis <sup>†</sup>	Piperacillin-tazobactam				Perforated appendicitis	5 days (after appropriate source control)
*Empiric anti-enterococcal therapy is recommended for patients with healthcare-associated intra-abdominal infections, those who have previously received cephalosporins, immunocompromised patients, and those with valvular heart disease or prosthetic intravascular materials. <sup>†</sup> <a href="#">LPCH Appendicitis Algorithm</a>								
<b>Traveler's diarrhea</b>	<i>E. coli</i> <i>Campylobacter</i> <i>Salmonella</i> , <i>Shigella</i> <i>Vibrio cholera</i> <i>Entamoeba histolytica</i> <i>Giardia</i> , <i>Blastocystis</i> <i>Cyclospora</i> , <i>Cystoisospora</i> <i>Cryptosporidium</i>	<b>First-line:</b> Azithromycin <b>Second-line:</b> Ciprofloxacin (should not be used for traveler's returning from SE Asia due to high prevalence of quinolone-resistant organisms)  <b>*Infants &lt;3 months old, severe disease, or CNS involvement:</b> Ceftriaxone <b>*Pediatric ID consult recommended</b>		Azithromycin: 10 mg/kg (max, 500 mg) PO daily X 3 days  Ciprofloxacin: 15 mg/kg (max, 500 mg) PO BID X 3 days  Ceftriaxone: 50 mg/kg IV q12h (max, 2000 mg) <b>*Duration dependent on severity of disease</b>		(37), (38)		
<b>Urinary tract infection (UTI)</b>	<i>E. coli</i> Other gram-negative bacteria ( <i>Klebsiella</i> , <i>Enterobacter</i> , <i>Citrobacter</i> ) <i>Enterococcus spp.</i>  NOTE: Ceftriaxone susceptibility is NOT a surrogate for cefazolin or cefdinir susceptibility	<u>Cystitis:</u> Nitrofurantoin OR Trimethoprim (TMP)-sulfamethoxazole (SMX)* <u>OR</u> Cephalexin*  <u>Pyelonephritis:</u> PO: Cephalexin*; if beta-lactam allergy, TMP-SMX*; IV: Ceftriaxone*  <b>*Enterococcus spp. are NOT susceptible to cephalosporins or TMP-SMX; usual drug of choice for Enterococcus spp. is Ampicillin (IV) OR Amoxicillin (PO).</b>  Ceftriaxone-resistant <i>E. coli</i> UTI may be treated with antibiotics other than meropenem, such as TMP-SMX PO, Ciprofloxacin or Levofloxacin, Gentamicin IV, Nitrofurantoin PO (if cystitis), and alternatively Amoxicillin-clavulanate PO (if cystitis) (see <a href="#">link</a> ).		<u>Cystitis</u> Nitrofurantoin X 5 days Trimethoprim-sulfamethoxazole X 3 days Cephalexin X 5 - 7 days  <u>Pyelonephritis:</u> 7 days of therapy is sufficient for children >6 months of age with uncomplicated pyelonephritis who improve within 3 days of starting antibiotics. Complicated pyelonephritis should be treated for 10 days.		(39-41)		
<b>Sexually transmitted infections</b>	Chancroid	< 45 kg: Azithromycin 20 mg/kg (max 1g) PO once OR ≥ 45 kg: Azithromycin 1 g PO once OR Ceftriaxone 250 mg IM once OR Ciprofloxacin 500 mg PO BID X 3 days				(42), (43)		

Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.
<a href="#">Sexually transmitted infections (cont.)</a>	Chlamydia	<p>Infants &lt;3 month of age: Erythromycin base ethylsuccinate 12.5 mg/kg PO QID X 14 days</p> <p>Children &lt;8 years old: Doxycycline 2 mg/kg (max 100 mg) PO BID X 7 days; Alternative: Azithromycin 20 mg/kg (max 1 g) PO as a single dose</p> <p>Children ≥8 years old, adolescents, and adults: Doxycycline* 2 mg/kg (max 100 mg) PO BID X 7 days Alternative: Azithromycin 20 mg/kg (max 1 g) PO once or Levofloxacin 500 mg PO daily X 7 days</p> <p>* Note that Doxycycline is contraindicated in the second and third trimester of pregnancy.</p>		
	Genital herpes	<p><u>1<sup>st</sup> episode:</u> PO: Valacyclovir 20 mg/kg (max 1 g) PO BID X 7-10 days OR Acyclovir 14 – 27 mg/kg (max 400 mg/dose) PO TID X 7-10 days IV: Acyclovir 5 - 10 mg/kg Q8h, change to PO when appropriate to complete therapy</p> <p><u>Episodic therapy:</u> Children and Adolescent: Acyclovir 20 mg/kg (max 400 mg) PO TID X 5 days OR Valacyclovir 20 mg/kg (max 500 mg) PO BID X 3 days OR Valacyclovir 20 mg/kg (max 1 g) PO daily X 5 days If adolescent or adult and HIV negative, can use high-dose acyclovir: Acyclovir 800 mg PO BID X 5 days OR Acyclovir 800 mg PO TID X 2 days</p> <p><u>Suppressive therapy:</u> Valacyclovir 10 mg/kg (max 500 mg) PO BID* OR Valacyclovir 20 mg/kg (max 1 g) PO once daily OR Acyclovir 20-25 mg/kg (max 400 mg) PO BID; *May be less effective for patients with very frequent recurrences (≥10 episodes per year)</p>		
	Gonorrhea	<p>Infants &amp; Children ≤45 kg with uncomplicated gonococcal vulvovaginitis, cervicitis, urethritis, pharyngitis, or proctitis: Ceftriaxone 50 mg/kg (max 500 mg) IM once</p> <p>Children &gt;45 kg, Adolescents and Adults with uncomplicated gonococcal vulvovaginitis, cervicitis, urethritis, pharyngitis, or proctitis: Ceftriaxone 500 mg IM once (if ≥150 kg, 1g) and if concurrent chlamydial infection has not been excluded, add Doxycycline 100 mg BID x 7 days (Alternative: Azithromycin 1g PO once) If beta-lactam allergy, Gentamicin 240 mg IM once + Azithromycin 2 g PO once</p>		
	Lymphogranuloma venereum	Doxycycline 100 mg PO BID X 21 days		
	Pelvic inflammatory disease (PID): <i>N. gonorrhoeae</i> <i>C. trachomatis</i> Vaginal flora (e.g., anaerobes, <i>G. vaginalis</i> , <i>H. influenzae</i> , enteric Gram-negative rods, GBS) <i>M. hominis</i> <i>U. urealyticum</i> <i>M. genitalium</i>	<p>Parenteral regimens: Cefoxitin 40 mg/kg (max 2 g) IV q6hr <b>AND</b> Doxycycline 2 mg/kg (max 100 mg) PO/IV q12hr or Clindamycin 13 mg/kg (max 900 mg) IV q8hr <b>AND</b> Gentamicin 2 mg/kg IV X 1, followed by 1.5 mg/kg IV q8hr</p> <p>Oral/IM regimens: Mild-to-moderately severe acute PID Ceftriaxone 50 mg/kg (max 500 mg; if ≥150 kg, 1g) IM once <b>AND</b> Doxycycline 2mg/kg (max 100 mg) PO BID X 14 days <b>AND</b> Metronidazole 15 mg/kg (max 500 mg) PO BID X 14 days or Cefoxitin 40 mg/kg (max 2 g) IM once <b>AND</b> Probenecid 1 g PO once <b>AND</b> Doxycycline 2 mg/kg (max 100 mg) PO BID X for 14 days <b>AND</b> Metronidazole 15 mg/kg (max 500 mg) PO BID X 14 days</p>		

Diagnosis		Usual Etiology	Suggested Empiric Therapy		Usual Duration of Treatment	Ref.
<a href="#">Sexually transmitted infections (cont.)</a>		Pubic lice	Permethrin 1% cream rinse or ivermectin 1% lotion; applied (up to 1 tube) to affected areas and washed off after 10 min OR Malathion 0.5% lotion applied to affected areas and washed off after 8–12h OR Ivermectin 250 mcg/kg PO once, then repeated 2 weeks later Note that Malathion lotion and Ivermectin oral are not FDA approved for treatment of public lice			
		Syphilis	<u>Congenital syphilis</u> *Pediatric ID consult recommended  <u>For Adolescents and Adults</u> <b>Primary, Secondary, and early latent (latent &lt;1 year) syphilis:</b> Benzathine penicillin G 2.4 million units/dose as a single dose intramuscularly  If treatment failure at 6-12 months, Benzathine penicillin G 50,000 units/kg IM weekly for three weeks (max 2.4 million units/dose)  For adolescents and adults (if infant or child, please contact Peds ID): <b>Late latent (latent &gt; 1 year), latent of unknown duration, late cardiovascular, gumma:</b> Benzathine penicillin G 2.4 million units IM X 3 weeks (consider EMLA before IM shot)  <u>Neurosyphilis</u> (children beyond neonatal period and adolescents) Penicillin G 50,000 units/kg q4-6hr Intravenously only (max 24 million units/day) X 10-14 days *If Penicillin allergic, consult ID			(42)
		Trichomoniasis	Children <45 kg: Metronidazole 15 mg/kg PO TID x 7 days (max: 2 g/day) Children ≥45kg and Adolescents: Metronidazole 2 g PO once OR Metronidazole 500 mg PO BID X 7 days			
		Vaginosis, bacterial	Metronidazole 15 mg/kg (max 500 mg) PO BID X 7 days <b>OR</b> Metronidazole gel 0.75%, one applicator (5 g) intravaginally once daily X 5 days <b>OR</b> Clindamycin cream 2%, one applicator (5 g) intravaginally at bedtime X 7days			
<b>Skin, Soft Tissues, and Skeletal System</b>						
<b>Bites, animal or human</b>		<i>S. aureus</i> Beta hemolytic streptococci (Group A) <i>Capnocytophaga canimorsus</i> <i>Pasteurella</i> (animal) <i>Eikenella</i> (human, animal)	1 <sup>st</sup> line: Amoxicillin-clavulanate (see <a href="#">link</a> )  2 <sup>nd</sup> line: Trimethoprim-sulfamethoxazole + Clindamycin  IV: Ampicillin-sulbactam If Penicillin allergy, Ceftriaxone + Clindamycin OR Trimethoprim-sulfamethoxazole + Clindamycin		Prophylaxis of high-risk bite wound: 3-5 days Cellulitis present: 10-14 days Tenosynovitis: 3 weeks Septic arthritis: 4 weeks Osteomyelitis: 6 weeks	(44), (45)
<b>Bone and joint infections (osteomyelitis, septic arthritis)</b>	Uncomplicated	<i>S. aureus</i> <i>S. pyogenes</i> <i>S. pneumoniae</i> <i>K. kingae</i>  Sickle cell disease: <i>Salmonella spp.</i>	Uncomplicated	Cefazolin If penicillin allergy, Clindamycin	Osteomyelitis: 4-6 weeks* Septic arthritis: 2-3 weeks*^  *Consider enteral transition once CRP down trending, afebrile ≥24hr, negative blood cultures ≥ 48hr, and improved use of affected bone or joint	(46-51)
			Complicated/ill-appearing/unimmunized	Ceftriaxone + Vancomycin		

Diagnosis		Usual Etiology	Suggested Empiric Therapy		Usual Duration of Treatment	Ref.
<b>Bone and joint infections (osteomyelitis, septic arthritis) (cont.)</b>	Complicated bone and joint infections	<i>S. aureus</i> <i>S. pyogenes</i> <i>S. pneumoniae</i> <i>Kingella spp.</i> (≤4 years) Enterobacterales <i>Pseudomonas spp.</i> Anaerobes  Infants (<3mo): GBS, <i>N. gonorrhoea</i>  Sickle cell disease: <i>Salmonella spp.</i>	Sickle cell disease	Ceftriaxone If beta-lactam allergy, Ciprofloxacin + Vancomycin	Drainage/debridement recommended  ^Durations as short as 10-14 days may be considered after 2-4 days of IV therapy in patients with <u>all</u> of the following: <ul style="list-style-type: none"> <li>• Infection due to <i>S. aureus</i>, <i>S. pyogenes</i>, <i>S. pneumoniae</i>, or <i>H. influenzae</i> type b</li> <li>• No adjacent osteomyelitis</li> <li>• Rapid clinical improvement</li> <li>• Consistent, progressive decrease in CRP by the end of the first week of treatment</li> </ul>	
			Atypical (immunocompromised, infants <3mo, duration of fever and/or pain >48-72hr, involves axial skeleton, presence of hardware)	Discuss with Pediatric ID consult service before start of empiric antibiotic		
			*Pediatric ID consult recommended			
<b>Cellulitis</b>	Group A streptococci (non-purulent) <i>S. aureus</i> (purulent)	<b>Non-purulent</b>		Cellulitis: 5 days Abscess: 5-10 days (Tailor duration based on resolution of signs and symptoms)	(8), (44)	
		<b>Mild</b>	Oral: Cephalexin If beta-lactam allergic, Clindamycin			
		<b>Moderate</b>	IV: Cefazolin If beta-lactam allergic, Clindamycin			
		<b>Severe</b>	Vancomycin + Piperacillin-tazobactam			
		<b>Purulent/abscess</b>				
		<b>Moderate*</b>	Trimethoprim-sulfamethoxazole OR Doxycycline OR Clindamycin			
		<b>Severe*</b>	Vancomycin			
		<b>Neonate</b>				
			*Pediatric ID consult recommended			
			*Incision and drainage recommended			
<b>Lymphadenitis</b>	Group A streptococci <i>S. aureus</i>	<u>Acute unilateral cervical lymphadenitis and moderate symptoms</u> (e.g., fever, ill-appearing, warm and/or tender adenitis) Cephalexin or Amoxicillin-clavulanate (see <a href="#">link</a> )  If concern for MRSA, Trimethoprim-sulfamethoxazole or Doxycycline can be considered		7 – 14 days depending on clinical response	(52)	
<b>Necrotizing fasciitis</b>	Beta hemolytic streptococci (Group A) <i>S. aureus</i> <i>V. vulnificus</i> <i>A. hydrophila</i> Peptostreptococci Polymicrobial (mixed aerobic-anaerobic microbes)	Vancomycin + Piperacillin-tazobactam + Clindamycin OR Piperacillin-tazobactam + Linezolid  <i>Prompt debridement required</i>  *Pediatric ID consult recommended		Depends on clinical response; duration usually prolonged (4-6 weeks)	(44), (53)	

Diagnosis	Usual Etiology	Suggested Empiric Therapy	Usual Duration of Treatment	Ref.
<b>Miscellaneous systemic infections</b>				
<a href="#">Febrile neutropenia</a>	<i>Pseudomonas</i> Enteric gram-negative bacilli <i>Staphylococci spp.</i> <i>Streptococci spp.</i> Yeast	*Refer to the LPCH F&N Guidelines		(54), (55)
<a href="#">Lyme disease</a>	<i>B. burgdorferi</i>  *Pediatric ID consult recommended  *Doxycycline is safe in children <8 years of age as long as duration does not exceed 21 days. Doxycycline has not been well studied in children < 8 years for durations >21 days	Early localized, erythema migrans,	Doxycycline 4.4 mg/kg per day divided BID (max 200 mg/day) for 10 days OR Amoxicillin 50 mg/kg per day divided TID (max 1.5 g/day) for 14 days OR Cefuroxime 30 mg/kg per day divided BID (max 1g/day) for 14 days	(56), (57)
		Isolated Bell's Palsy	Doxycycline 4.4 mg/kg per day divided BID (max 200 mg/day) for 14 days. *Amoxicillin and cefuroxime have not been studied for the treatment of Lyme Bell's Palsy	
		Arthritis	<b>Initial episode:</b> As for early localized disease with a duration of 28 days	
			<b>Recurrent arthritis</b> despite adequate prior oral therapy: Repeat second course of oral antibiotics as above x4 weeks OR Ceftriaxone 50-75 mg/kg (max 2 g) IV once daily for 14-28 days	
		Carditis	Ceftriaxone 50-75 mg/kg once daily for 14-21 days. Can consider transitioning to Doxycycline once patient improving	
		Meningitis, neuroborreliosis	Ceftriaxone 50-75 mg/kg (max 2 g) IV once daily for 14 days OR Doxycycline 4.4 mg/kg per day divided BID (max 200 mg/day) for 14 days	

❖ Please visit the [Antimicrobial Stewardship Program \(ASP\)](#) website located in the Housestaff Manual for more useful resources, including:

- [LPCH antibiogram and antimicrobial dosing quick reference](#)
- [Antimicrobial monitoring](#)
- [Neonatal Intensive Care Unit \(NICU\) late onset sepsis algorithm](#)
- Vancomycin Dosing Guidelines:
  - [Vancomycin Guideline for Neonates and Children](#)
  - [Vancomycin Guideline in Hemodialysis](#)
  - [Initial Vancomycin Dosing Guideline for Surgical Patients up to 1 Week Post-Op](#)
- [Restricted antimicrobials and the appropriate approval process](#)
- [Palivizumab \(Synagis®\) guidelines](#)
- [Surgical prophylaxis guidelines](#)
- [Amoxicillin-clavulanate dosing and formulation selection guide](#)
- Other clinical practice guidelines and clinician education

## References:

1. Tunkel AR, Hartman BJ, Kaplan SL, et al. Practice guidelines for the management of bacterial meningitis. *Clin Infect Dis* **2004**; 39:1267-1284.
2. Ku LC, Boggess KA, Cohen-Wolkowicz M. Bacterial meningitis in infants. *Clinics in perinatology* **2015**; 42:29-45.
3. Bonfield CM, Sharma J, Dobson S. Pediatric intracranial abscesses. *J Infect.* **2015** Jun;71 Suppl 1:S42-6. doi: 10.1016/j.jinf.2015.04.012. Epub 2015 Apr 24. PMID: 25917804
4. Tunkel AR, et al. 2017 Infectious Diseases Society of America's clinical practice guidelines for healthcare-associated ventriculitis and meningitis. *Clin Infect Dis* **64.6** **2017**; 64(6): e34-e65.
5. Lieberthal AS, Carroll AE, Chonmaitree T, et al. The diagnosis and management of acute otitis media. *Pediatrics* **2013**; 131:e964-e999.
6. Neff L, Newland JG, Sykes KJ, Selvarangan R, Wei JL. Microbiology and antimicrobial treatment of pediatric cervical lymphadenitis requiring surgical intervention. *Int J Pediatr Otorhinolaryngol.* **2013** May;77(5):817-20.
7. Weinstock MS, Patel NA, Smith LP. "Pediatric cervical lymphadenopathy." **2018**;39 (9):433-443.
8. Rybak MJ, Le J, Lodise TP, et al. Therapeutic Monitoring of Vancomycin for Serious Methicillin-resistant Staphylococcus aureus Infections: A Revised Consensus Guideline and Review by the American Society of Health-system Pharmacists, the Infectious Diseases Society of America, the Pediatric Infectious Diseases Society, and the Society of Infectious Diseases Pharmacists. *Clin Infect Dis.* **2020** Sep 12;71(6):1361-1364.
9. Griffith D.E., Aksamit T., Brown-Elliott B.A., Catanzaro A., et al. An official ATS/IDSA statement: diagnosis, treatment, and prevention of nontuberculous mycobacterial diseases. *Am J Respir Crit Care Med.* **2007**;175(4):367-416.
10. Nahid P, Dorman SE, Apilana N. Official American Thoracic Society/Centers for Disease Control and Prevention/Infectious Diseases Society of America Clinical Practice Guidelines: treatment of drug-susceptible tuberculosis. *Clin Infect Dis.* **2016**;63(7):e147-9.
11. Nayak JL et al: Epiglottitis. In: Bennett JE et al, eds: Mandell, Douglas, and Bennett's Principles and Practice of Infectious Disease, Updated Edition. 8th ed. Philadelphia, PA: Saunders; 2015:785-8
12. Lichtor JL, et al. Epiglottitis It Hasn't Gone Away. *Anesthesiology: The Journal of the American Society of Anesthesiologists* **2016**;24.6: 1404-1407.
13. Hollis S, Evans K. Management of malignant (necrotizing) otitis externa. *The Journal of Laryngology & Otology* **2011**;125:1212-1217.
14. Mion M, Bovo R, Marchese-Ragona R, et al. Outcome predictors of treatment effectiveness for fungal malignant external otitis: a systematic review. *Acta Otorhinolaryngol Ital* **2015**; 35:307.
15. Dietz E, Mangus C. Microbiology and infectious disease. In: Engorn B, Flerlage J, eds. The Harriet Lane Handbook. 20<sup>th</sup> ed. Philadelphia, PA: Elsevier Inc; 2015: 380-414.
16. Carter L, Callon W. Mastoiditis. *Pediatr Rev.* 2024 Sep 1;45(9):540-542. doi: 10.1542/pir.2023-006249. PMID: 39217116.
17. World Health Organization. Chronic suppurative otitis media – Burden of illness and management options. Geneva, Switzerland. **2004**
18. Caviglia I, Techera A, García G. Antimicrobial therapies for odontogenic infections in children and adolescents. *Odontoestomatología* **2016**;18.27: 4-15.
19. Kaneko A, et al. The 2016 JAID/JSC guidelines for clinical management of infectious disease– Odontogenic infections. *J Infect Chemother.* **2018**;24(5):320-324.
20. Holmes CJ, Pellecchia R. Antimicrobial Therapy in Management of Odontogenic Infections in General Dentistry. *Dent. Clin. N. Am.* **2016**;60:497-507.
21. Apostolopoulos NJ, Nikolopoulos TP, Bairamis TN. Peritonsillar abscess in children. Is incision and drainage an effective management? *Int J Pediatr Otorhinolaryngol* **1995**; 31:129.
22. American Academy of Pediatrics. Pertussis (Whooping Cough). In: Kimberlin DW, Brady MT, Jackson MA, Long SS, eds. *Red Book: 2018 Report of the Committee on Infectious Diseases.* American Academy of Pediatrics; **2018**; 620-634.
23. Shulman ST, Bisno AL, Clegg HW, et al. Clinical practice guideline for the diagnosis and management of group A streptococcal pharyngitis: 2012 update by the Infectious Diseases Society of America. *Clin Infect Dis* **2012**; 55(10):1279-1282.
24. Metlay JP, Waterer GW, Long AC, Anzueto A, Brozek J, Crothers K, et al. Diagnosis and Treatment of Adults with Community-acquired Pneumonia. An Official Clinical Practice Guideline of the American Thoracic Society and Infectious Diseases Society of America. *Am J Respir Crit Care Med.* **2019**;200(7):e45-e67.
25. El-Solh AA, Pietrantoni C, Bhat A, Aquilina AT, Okada M, Grover V, Gifford N. Microbiology of severe aspiration pneumonia in institutionalized elderly. *Am J Respir Crit Care Med.* **2003** Jun 15;167(12):1650-4.
26. Bradley JS, Byington CL, Shah SS, et al. The management of community-acquired pneumonia in infants and children older than 3 months of age: clinical practice guidelines by the Pediatric Infectious Diseases Society and the Infectious Diseases Society of America. *Clin Infect Dis* **2011**; 53(7):e25:76.
27. American Thoracic Society and Infectious Diseases Society of American. Guidelines for the management of adults with hospital-acquired, ventilator-associated, and healthcare-associated pneumonia. *Am J Respir Care Med* **2005**; 171(4):388-416.
28. Chow AW, Benninger MS, Brook I, et al. IDSA clinical practice guideline for acute bacterial rhinosinusitis in children and adults. *Clin Infect Dis* **2012**; 54(8):e72-112.
29. Wald ER, et al. Clinical practice guideline for the diagnosis and management of acute bacterial sinusitis in children aged 1 to 18 years. *Pediatrics* **2013**: e262-e280.

30. Starkey CR, Steele RW. Medical management of orbital cellulitis. *Pediatr Infect Dis J* **2001**; 20:1002.
31. Howe L, Jones NS. Guidelines for the management of periorbital cellulitis/abscess. *Clin Otolaryngol Allied Sci.* **2004** Dec;29(6):725-8. doi: 10.1111/j.1365-2273.2004.00889.x.
32. Baltimore RS, Gewitz M, Baddour LM, et al. Infective endocarditis in childhood: 2015 update: a scientific statement from the AHA. *Circulation* **2015**; 132(15):1487-1515.
33. Chong YP, Moon SM, Bang KM, et al. Treatment duration for uncomplicated Staphylococcus aureus bacteremia to prevent relapse: analysis of a prospective observational cohort study. *Antimicrob Agents Chemother* **2013**; 57(3):1150-1156.
34. Duguid RC, Al Reesi M, Bartlett AW, Palasanthiran P, McMullan BJ. Impact of Infectious Diseases Consultation on Management and Outcome of Staphylococcus aureus Bacteremia in Children. *J Pediatric Infect Dis Soc.* **2021** May 28;10(5):569-575. doi: 10.1093/jpids/piaa155.
35. Johnson S, Lavergne V, Skinner AM, et al. Clinical Practice Guideline by the Infectious Diseases Society of America (IDSA) and Society for Healthcare Epidemiology of America (SHEA): 2021 Focused Update Guidelines on Management of *Clostridioides difficile* Infection in Adults. *Clinical Infectious Diseases.* 2021; 73:e1029-e44.
36. Solomkin JS, Mazuski JE, Bradley JS, et al. Diagnosis and management of complicated intra-abdominal infection in adults and children: guidelines by the Surgical Infection Society and the Infectious Diseases Society of America. *Clin Infect Dis* **2010**; 50:133-164.
37. Bradley JS and Nelson JD. Nelson's Pediatric Antimicrobial Therapy. American Academy of Pediatrics. Elk Grove Village, IL 2015. P.75.
38. Shane, Andi L., et al. 2017 Infectious Diseases Society of America clinical practice guidelines for the diagnosis and management of infectious diarrhea. *Clinical Infectious Diseases* 65.12 **2017**: e45-e80.
39. American Academy of Pediatrics, Committee on Quality Improvement and Management, Subcommittee on Urinary Tract Infection. Urinary tract infection: clinical practice guideline for the diagnosis and management of the initial UTI in febrile infants and children 2-24 months. *Pediatrics* **2011**; 128(3):595-610.
40. Wang ME, Lee V, Greenhow TL, Beck J, Bendel-Stenzel M, Hames N, McDaniel CE, King EE, Sherry W, Parmar D, Patrizi ST, Srinivas N, Schroeder AR. Clinical Response to Discordant Therapy in Third-Generation Cephalosporin-Resistant UTIs. *Pediatrics.* **2020** Feb;145(2):e20191608. doi: 10.1542/peds.2019-1608.
41. Fox MT, et al. Comparative effectiveness of antibiotic treatment duration in children with pyelonephritis. *JAMA Network Open.* **2020**;3.5:e203951-e203951.
42. Workowski K, et al. Sexually transmitted diseases treatment guidelines, 2021. *MMWR Recomm Rep.* **2021**; 70(4):1-192.
43. St. Cyr S, Barbee L, Workowski KA, et al. Update to CDC's Treatment Guidelines for Gonococcal Infection, 2020. *MMWR Morb Mortal Wkly Rep* 2020;69:1911–1916.
44. Stevens DL, Bisno AL, Chambers HF, et al. Practice guideline for the diagnosis and management of skin and soft tissue infections: 2014 update by the Infectious Diseases Society of America. *Clin Infect Dis* **2014**; 59(2):147-159.
45. Patil PD, Panchabhai TS, Galwankar SC. Managing human bites. *Journal of Emergencies, Trauma and Shock.* **2009**;2.3:186.
46. Peltola H, Paakkonen M. Acute osteomyelitis in children. *N Engl J Med* **2014**;370(4):352-360.
47. Hatzenbuehler J, Pulling TJ. Diagnosis and management of osteomyelitis. *Am Fam Physician.* **2011.** 84(9): 1027-1033.
48. Shariff KA, Richards EP, Townes JM. Clinical management of septic arthritis. *Curr Rheumatol Rep.* **2013.** 15: 332. 1-9
49. Horowitz DL, Katzap E. Approach to septic arthritis. *Am Fam Physician.* **2011.** 84(6): 653-660.
50. Peltola H, Paakkonen M, Kallio P, et al. Prospective, randomized trial of 10 days versus 30 days of antimicrobial treatment, including a short-term course of parenteral therapy, for childhood septic arthritis. *Clin Infect Dis* **2009**; 48(9):1201-1210.
51. Guideline on Diagnosis and Management of Acute Bacterial Arthritis in Pediatrics [published correction appears in *J Pediatric Infect Dis Soc.* 2024 Jul 13; piae065. doi: 10.1093/jpids/piae065]. *J Pediatric Infect Dis Soc.* 2024;13(1):1-59. doi:10.1093/jpids/piad089
52. Pecora F, Abate L, Scavone S, Petrucci I, Costa F, Caminiti C, Argentiero A, Esposito S. Management of Infectious Lymphadenitis in Children. *Children (Basel).* **2021** Sep 27;8(10):860. doi: 10.3390/children8100860.
53. Misiakos EP, Bagias G, Patapis P, et al. Current concepts in the management of necrotizing fasciitis. *Front Surg* **2014**; 1:36. doi: 10.3389/fsurg.2014.00036. eCollection 2014.
54. Freifeld AG, Bow EJ, Spkowitz KA, et al. Clinical Practice Guideline for the Use of Antimicrobial Agents in Neutropenic Patients with Cancer: 2010 Update by the Infectious Diseases Society of America. *Clin Infect Dis* **2011**; 52(4):e56-e93.
55. Lehrnbecher T, et al. Guideline for the management of fever and neutropenia in children with cancer and hematopoietic stem-cell transplantation recipients: 2017 update. *J Clin Oncol.* **2017**;35(18):2082-2094.
56. Lantos PM, Rumbaugh J, Bockenstedt LK, et al. Clinical Practice Guidelines by the Infectious Diseases Society of America (IDSA), American Academy of Neurology (AAN), and American College of Rheumatology (ACR): 2020 Guidelines for the Prevention, Diagnosis, and Treatment of Lyme Disease. *Arthritis Rheumatol.* 2021;73(1):12-20. doi:10.1002/art.41562
57. American Academy of Pediatrics Report of the Committee on Infectious Diseases. Red Book. 32<sup>nd</sup> edition. 2021-2024.

58. Maraolo AE, Mazzitelli M, Zappulo E, et al. Oral Vancomycin Prophylaxis for Primary and Secondary Prevention of *Clostridioides difficile* Infection in Patients Treated with Systemic Antibiotic Therapy: A Systematic Review, Meta-Analysis and Trial Sequential Analysis. *Antibiotics (Basel)*. 2022;11(2):183. Published 2022 Jan 30. doi:10.3390/antibiotics11020183
59. Bao H, Lighter J, Dubrovskaya Y, et al. Oral Vancomycin as Secondary Prophylaxis for *Clostridioides difficile* Infection. *Pediatrics*. 2021;148(2):e2020031807. doi:10.1542/peds.2020-031807
60. Woods CR, Bradley JS, Chatterjee A, et al. Clinical Practice Guideline by the Pediatric Infectious Diseases Society (PIDS) and the Infectious Diseases Society of America (IDSA): 2023
60. Wolf J, Kalocsai K, Fortuny C, et al. Safety and efficacy of fidaxomicin and vancomycin in children and adolescents with clostridioides (clostridium) difficile infection: a phase 3, multicenter, randomized, single-blind clinical trial (SUNSHINE). *Clin Infect Dis*. 2020;71(10):2581–2588. doi: 10.1093/cid/ciz1149
61. Alsoubani M, Chow JK, Rodday AM, Kent D, Snyderman DR. Comparative Effectiveness of Fidaxomicin vs Vancomycin in Populations With Immunocompromising Conditions for the Treatment of *Clostridioides difficile* Infection: A Single-Center Study. *Open Forum Infect Dis*. 2023;11(1):ofad622. Published 2023 Dec 8. doi:10.1093/ofid/ofad622
62. Zhao Z, Wu Y, Geng X, Yuan C, Fu Y, Yang G. Efficacy of fidaxomicin versus vancomycin in the treatment of Clostridium difficile infection: A systematic meta-analysis. *Medicine (Baltimore)*. 2024;103(32):e39213. doi:10.1097/MD.00000000000039213
63. Rao K, Zhao Q, Bell J, et al. An Open-Label, Randomized Trial Comparing Fidaxomicin With Oral Vancomycin for the Treatment of Clostridioides difficile Infection in Hospitalized Patients Receiving Concomitant Antibiotics for Concurrent Infections. *Clin Infect Dis*. 2024;78(2):277-282. doi:10.1093/cid/ciad606
64. Dop D, Marcu IR, Padureanu V, et al. Clostridium difficile infection in pediatric patients (Review). *Biomed Rep*. 2023;20(2):18. Published 2023 Dec 8. doi:10.3892/br.2023.1706
65. American Academy of Pediatrics Committee on Infectious Diseases (2024). Principles of appropriate use of antimicrobial agents. In D.W. Kimberlin, M.T. Brady, M.A. Jackson, & S.S. Long (Eds.), *Red Book: 2024-2027 Report of the Committee on Infectious Diseases* (33<sup>rd</sup> ed.). American Academy of Pediatrics. [https://doi.org/10.1542/9781610027373-S4\\_003\\_006](https://doi.org/10.1542/9781610027373-S4_003_006)
- Falagas ME, Karageorgopoulos DE, Grammatikos AP, Matthaiou DK. Effectiveness and safety of short vs. long duration of antibiotic therapy for acute bacterial sinusitis: a meta-analysis of randomized trials. *Br J Clin Pharmacol*. 2009;67(2):161-171. doi:10.1111/j.1365-2125.2008.03306.x