



Pediatric Skin and Soft Tissue Infection Treatment Guideline

Disclaimer: this guideline should not replace clinical judgement.

Degree of Severity Definition		
Severity	Purulent	Non-Purulent
Mild	Incision and drainage (I&D) only	
Moderate	Systemic signs of infection	Cellulitis/erysipelas with no focus of purulence
Severe	Failed I&D Sepsis Severely Immunocompromised	Sepsis Severely Immunocompromised Presence of bullae or sloughing

Management of <u>non-purulent SSTI</u>				
Type of Infection	Organisms	Preferred Treatment	Alternative Treatment (PCN allergy)	Duration
Cellulitis and Erysipelas *Erysipelas has defined borders *Blood cultures, tissue aspirates, and skin biopsies are NOT routinely recommended due to low yield	Beta-hemolytic Streptococci: Group A – S. <i>pyogenes</i> (most common), Group B – <i>S. agalactiae</i> , Groups C, G, F <i>Staphylococcus aureus</i> only if: large open wound, IV drug user, penetrating trauma, active <i>S. aureus</i> infection at another site	Mild Treatment Options (oral): Penicillin VK 20 mg/kg/dose PO q8h (max 500 mg/dose) Amoxicillin 12.5 mg/kg/dose PO q8h (max 500 mg/dose)	Cephalexin 12.5 mg/kg/dose PO q6h (max 500 mg/dose) Clindamycin 5-10 mg/kg/dose PO q8h (max 450 mg/dose)	5 days
		Moderate Treatment Options (intravenous): Penicillin G 60,000 to 100,000 units/kg/dose IV q6h (max 4 million units/dose) Ampicillin 25 mg/kg IV q6h (max 2000 mg/dose)	Cefazolin 33 mg/kg IV q8h (max 1000 mg/dose) Clindamycin 10 mg/kg/dose IV q8h (max 600 mg/dose)	Duration is not contingent on erythema resolution alone, may extend to 7-10 days if slow clinical improvement
		Severe Treatment Options (intravenous antibiotics): See Necrotizing Fasciitis recommendations below Consider ID consult		



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Impetigo * Oral antimicrobials recommended for numerous lesions or outbreaks affecting several people to decrease transmission	<i>S. aureus</i> <i>S. pyogenes</i>	Mild: Mupirocin (topical) twice daily Moderate-Severe: <u>Empiric/MSSA</u> Cephalexin 12.5 mg/kg PO q6h (max 500 mg/dose) <u>MRSA suspected/confirmed.</u> Clindamycin 5-10 mg/kg PO q8h (max 450 mg/dose)	Moderate-Severe: <u>Empiric/MSSA</u> Clindamycin 10 mg/kg PO q6h (max 600 mg/dose) <u>MRSA</u> SMX/TMP 5 mg TMP/kg PO q12h (max 160 mg TMP/dose) Doxycycline* 2 mg/kg PO q12h (max 100 mg/dose)	Mupirocin: 5 days All others: 7 days	
Folliculitis	<i>S. aureus</i> <i>Pseudomonas aeruginosa</i> (hot tubs)		No antimicrobials Warm compresses Gentle cleanser		
Necrotizing Fasciitis	Empirically: Broad spectrum gram positive, gram negative and anaerobic coverage Most common organisms: Group A Streptococci (GAS) Gas Gangrene: <i>Clostridium perfringens</i> , <i>Clostridium septicum</i>	Emergent surgical consultation	Empiric: Vancomycin 15 mg/kg IV q6h (max 2000 mg/dose)** PLUS Piperacillin/tazobactam 75 mg pip/kg/dose IV q6h (max 3.375 g/dose) Identified GAS or <i>Clostridium</i>: Penicillin 60,000 to 100,000 units/kg/dose IV q6h (max 4 million units/dose) PLUS Clindamycin 10 mg/kg IV q6h (max 900 mg/dose)	Empiric: Linezolid <12 years: 10 mg/kg IV/PO q8h (max 600 mg/dose) ≥12 years: 600 mg IV/PO q12h PLUS Cefepime 50 mg/kg IV q8h (max 2000 mg/dose) PLUS Metronidazole 10 mg/kg IV/PO q6h (max 500 mg/dose) Meropenem 20 mg/kg IV q8h (max 1000 mg/dose)	Dependent upon surgical debridement/source control *Clindamycin used in combination for toxin binding should be discontinued after 48-72 hours

*Doxycycline is NOT recommended for children < 8 years old but may be used in life-threatening situations

** Vancomycin may be adjusted per protocol based on age and renal function

*** Addition of clindamycin has been shown to reduce the in vitro release of streptococcal pyrogenic exotoxins, however there is still a lack of clinical prospective trials strongly recommending its use in this setting. **Addition is not needed if linezolid is part of the empiric regimen as linezolid has been shown to reduce toxin production.** Surgical intervention remains the most important treatments to manage necrotic spread.



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Management of <u>purulent SSTI</u>				
Type of Infection	Organisms	Preferred Treatment	Alternative Treatment (IV, PCN allergy)	Duration
Abscesses (Furuncles, Carbuncles) *For moderate/severe, send purulent material for culture and sensitivity	<i>S. aureus</i>		Mild I&D only	
		Moderate/Severe I&D plus systemic antibiotics Empiric/MRSA: Clindamycin 10 mg/kg IV/PO q8h (max 600 mg/dose) MSSA: Cephalexin 12.5 mg/kg PO q6h (max 500 mg/dose) Cefazolin 33 mg/kg IV q8h (max 1000 mg/dose)	Empiric/MRSA: Vancomycin 15 mg/kg IV q6h (max 2000 mg/dose)** SMX/TMP 5 mg TMP/kg IV/PO q12h (max 160 mg TMP/dose) Linezolid <12 years: 10 mg/kg IV/PO q8h (max 600 mg/dose) ≥12 years: 600 mg IV/PO q12h MSSA: Nafcillin 25 mg/kg IV q6h (max 2000 mg/dose)	5 days May extend to 7-10 days if slow clinical improvement
Preseptal (Periorbital) Cellulitis	<i>S. aureus</i> <i>S. epidermidis</i> <i>S. pyogenes</i> <i>H. influenzae</i> (unimmunized)	Clindamycin 10 mg/kg IV/PO q6h (max 600 mg/dose) Rhinosinusitis/unimmunized: Amoxicillin/clavulanate (Formulation 7:1) 22.5 mg/kg PO q12h (max 875 mg amox/dose)	Rhinosinusitis/unimmunized: Cefdinir 7 mg/kg PO BID (max 300 mg/dose) OR Ceftriaxone 50-75 mg/kg IV q24h (max 2000 mg/dose)	5 days May extend to 7-10 days if slow clinical improvement



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Orbital Cellulitis	<p><i>S. aureus</i> <i>S. epidermidis</i> <i>S. pyogenes</i></p> <p>Other organisms: <i>S. anginosus</i> Neisseria spp <i>M. catarrhalis</i> Oral anaerobes <i>H. influenzae</i> (unimmunized)</p>	<p>Clindamycin 10 mg/kg IV q6h (max 600 mg/dose) PLUS Ceftriaxone 50-75 mg/kg IV q24h (max 2000 mg/dose)</p> <p>Suspected intracranial extension: Vancomycin 20 mg/kg IV q6h (max 2000 mg/dose)** PLUS Meropenem 20 mg/kg IV q8h (max 1000 mg/dose) OR Levofloxacin</p> <p><u>6 months-5 years:</u> 10 mg/kg IV q12h (max 375 mg/dose) <u>≥5 years:</u> 10 mg/kg IV q24h (max 750 mg/dose)</p>	<p>Suspected intracranial extension: Vancomycin 20 mg/kg IV q6h (max 2000 mg/dose)** PLUS Meropenem 20 mg/kg IV q8h (max 1000 mg/dose) OR Levofloxacin</p> <p><u>6 months-5 years:</u> 10 mg/kg IV q12h (max 375 mg/dose) <u>≥5 years:</u> 10 mg/kg IV q24h (max 750 mg/dose)</p>	Minimum 10-14 days, dependent on symptom resolution
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** Vancomycin may be adjusted per protocol based on age and renal function

Surgical Site Infections

Type of Infection	Suspected Organisms	Recommended Treatment Options
Surgical Site Infections *Culture and sensitivity for any purulent material *Short course (24-48 hours) recommended ONLY for patients with significant systemic response	<p><48 hours: <i>S. pyogenes</i>, Clostridium spp</p> <p>>48 hours: <i>S. aureus</i></p>	<p>Suture removal plus I&D</p> <p>MRSA Clindamycin 10 mg/kg IV/PO q6h (max 600 mg/dose) Vancomycin 15 mg/kg IV q6h (max 2000 mg/dose)** Linezolid <u><12 years:</u> 10 mg/kg IV/PO q8h (max 600 mg/dose) <u>≥12 years:</u> 600 mg IV/PO q12h</p> <p>MSSA Cefazolin 33 mg/kg IV q8h (max 2000 mg/dose) Nafcillin 25 mg/kg IV q6h (max 2000 mg/dose)</p>
	Consider adding coverage for gram negatives and anaerobes for surgeries involving: <ul style="list-style-type: none"> • GI tract • Perineum • Female genital tract 	<p>Ceftriaxone 50-75 mg/kg IV q24h (max 2000 mg/dose) PLUS Metronidazole 10 mg/kg IV/PO q8h (max 500 mg/dose)</p> <p>PCN allergy: Levofloxacin <u>6 months-5 years:</u> 10 mg/kg IV q12h (max 375 mg/dose) <u>≥5 years:</u> 10 mg/kg IV q24h (max 750 mg/dose) PLUS Metronidazole 10 mg/kg IV/PO q8h (max 500 mg/dose)</p>



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Bite Wounds					
Tetanus and Rabies vaccines as appropriate					
For patients with wounds that do not appear infected, consider 3-5 days of pre-emptive antimicrobial therapy for:					
IDSA Prophylaxis Recommendations			Red Book Prophylaxis Recommendations		
<ul style="list-style-type: none"> • Immunocompromised or asplenic • Advanced liver disease • Preexisting or resultant edema of the affected area • Moderate to severe injuries (especially hands or face) • Injuries involving periosteum or joint capsule 			<ul style="list-style-type: none"> • Moderate or severe bite wounds, especially if edema or crush injury is present • Puncture wounds, especially if penetration of bone, tendon sheath, or joint has occurred • Deep or surgically closed facial bite wounds • Hand and foot bite wounds • Genital area bite wounds • Wounds in immunocompromised and asplenic children • Wounds exhibiting signs of infection • Cat bite wounds 		
Source of bite	Common Organisms	Antimicrobial Agents			
		Oral	Oral, PCN allergy	IV	IV, PCN allergy
Dog, cat, other mammal	<i>Pasteurella</i> spp <i>S. aureus</i> Streptococci <i>Capnocytophaga</i> spp <i>Moraxella</i> spp <i>Neisseria</i> spp <i>Corynebacterium</i> spp Anaerobes	Amoxicillin/clavulanate (formulation 7:1) 12.5 mg/kg PO q12h (max 875 mg amox/dose)	Clindamycin 10 mg/kg IV/PO q6h (max 600 mg/dose) PLUS Cefuroxime 15 mg/kg BID (max 500 mg/dose) OR Cefdinir 7 mg/kg PO BID (max 300 mg/dose) OR TMP/SMX 5 mg TMP/kg IV/PO q12h (max 160 mg TMP/dose)	Ampicillin-sulbactam 25 mg ampicillin/kg IV q6h (max 2000 mg ampicillin/dose [3000 mg ampicillin-sulbactam])	
Human	Streptococci <i>S. aureus</i> <i>Eikenella corrodens</i> <i>Haemophilus</i> spp Anaerobes				Ceftriaxone 50-75 mg/kg IV q24h (max 2000 mg/dose) OR TMP/SMX 5 mg TMP/kg q12h (max 160 mg TMP/dose)
Reptile	Enteric gram-negatives Anaerobes			Ampicillin-sulbactam 25 mg ampicillin/kg IV q6h (max 2000 mg ampicillin/D Dose [3000 mg ampicillin-sulbactam]) PLUS Gentamicin * 5 mg/kg IV q24h	

*Gentamicin: pharmacy consult is recommended



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References

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