

MEDICAL CENTER

Vanderbilt Adult Antimicrobial Stewardship Program

VASP Diabetic Foot Infection & Diabetic Foot Osteomyelitis – Inpatient Management

This guidance document is meant to provide general recommendations and does not supersede clinical decision making. It does not include comprehensive recommendations for surgical management (including need for revascularization), wound care, offloading, smoking cessation, or glycemic control.

I. Foundational Principles

- a. Do **NOT** use antibiotics to treat uninfected wounds as patient harm outweighs benefit.
- b. **ALL** wounds are colonized with microorganisms and superficial wound cultures should not be collected nor used to make treatment decisions.
- c. Do **NOT** administer antibiotics to clinically stable patients until deep tissue or operative cultures are obtained.
- d. Failure of oral antibiotics used for < 48 hours is **NOT** a reason to use broad-spectrum antibiotics.

II. Diagnosis

- a. Based on presence of clinical signs or symptoms of local or systemic infection in patients with diabetes
 - i. Inflammation of any part of the foot
 - ii. Presence of ≥ 2 SIRS criteria
- b. Assess severity based on validated scoring metrics (see [Table 1](#)).
- c. Use of appropriate imaging modalities (e.g. X-ray or MRI) to determine extent of infection and/or bone involvement at discretion of treating physician.
- d. Use of inflammatory markers such as ESR or CRP at baseline may help determine level of inflammation with periodic trending (e.g. weekly) thereafter. Do **NOT** repeat these tests daily.
- e. Surgical management and wound care at discretion of treating physician.
- f. Appropriate types of cultures:
 - i. Deep tissue/wound culture after debridement
 - ii. Aseptically collected bone sample when osteomyelitis suspected
 - iii. Blood cultures only in patients with systemic findings (e.g. ≥ 2 SIRS criteria)
 - iv. Consider obtaining MRSA nasal PCR (ensure collected before nasal decolonization)*

III. MRSA and *P. aeruginosa* Risk Factors

- a. MRSA (≥ 1)
 - i. Previous MRSA infection or colonization
 - ii. Hospitalization within previous 90 days **AND** IV antibiotic use
 - iii. IV drug use
 - iv. Chronic hemodialysis
- b. *P. aeruginosa* (≥ 1)
 - i. Previous *P. aeruginosa* infection or colonization
 - ii. Water exposure or macerated wounds
 - iii. Hospitalization within previous 90 days **AND** IV antibiotic use

- iv. Immunocompromise (solid organ transplant, bone marrow transplant, active chemotherapy, chronic steroids, neutropenia, HIV with CD4 < 200)

IV. Antimicrobial dosing

- a. Reference available on the VASP website at the following [link](#).

*MRSA nasal PCR has ~90% negative predictive value for diabetic foot infections.

Table 1: Diabetic Foot Infection Severity Classification

Clinical Criteria	IWGD/IDSA Classification
No systemic or local symptoms or signs of infection	1/Uninfected
Infected (must have ≥ 2 of the following): <ul style="list-style-type: none"> • Local swelling or induration • Erythema > 0.5 but < 2 cm around the wound • Local tenderness or pain • Local increased warmth • Purulent discharge Rule out other causes of inflammation (e.g. trauma, gout, acute charcot foot, fracture, thrombosis, venous stasis)	2/Mild
Infected (no systemic findings): <ul style="list-style-type: none"> • Erythema ≥ 2 cm from wound margin (including lymphangitis, deep tissue abscess, gangrene), and/or • Deep tissue involvement (e.g. tendon, muscle, joint) 	3/Moderate
Bone Infection (osteomyelitis)	Add "O"
Infected (with systemic findings): <ul style="list-style-type: none"> • Any of the local manifestations above, AND • At least 2 SIRS criteria <ul style="list-style-type: none"> ○ Temperature > 38°C, ○ Heart rate > 90bpm, ○ Respiratory rate > 20bpm, ○ WBC > 12,000/mm³ 	4/Severe
Bone Infection (osteomyelitis)	Add "O"

V. Treatment Recommendations

Classification	Pathogens	Empiric Antibiotic Therapy*	Duration of Treatment
1/Uninfected	Do NOT administer antibiotics. Do NOT culture the wound.		
2/Mild	GPC (β -hemolytic streptococci and <i>S. aureus</i>)	<p>Preferred:</p> <ul style="list-style-type: none"> Cephalexin 500mg PO QID or 1g TID Cefadroxil 1g PO BID (upon discharge only) <p>Alternative:</p> <ul style="list-style-type: none"> Amoxicillin/clavulanate 875/125mg PO BID <p>MRSA only with risk factors:</p> <ul style="list-style-type: none"> Doxycycline 100mg PO BID (add to β-lactam) Trimethoprim/sulfamethoxazole 5-8mg/kg/day (alone)—Preferred if severe penicillin allergy 	7 – 14 days
3/Moderate	<p>GPC \pm GNR \pm anaerobes</p> <ul style="list-style-type: none"> Do NOT administer antibiotics to clinically stable patients until deep tissue or operative cultures are obtained. Cultures from the last 6 months may help empiric antibiotic selection. Adjust empiric antibiotic regimen based on deep culture and susceptibility results 	<p>Preferred:</p> <ul style="list-style-type: none"> Ampicillin/sulbactam 3g IV Q6H Ceftriaxone 2g IV daily \pm PO metronidazole 500mg Q12H <p><i>P. aeruginosa</i> only with risk factors (change to):</p> <ul style="list-style-type: none"> Piperacillin/tazobactam 4.5g IV Q8H Cefepime 2g IV Q8H \pm metronidazole 500mg PO Q12H Levofloxacin 750mg PO daily \pm metronidazole 500mg PO Q12H <p>MRSA only with risk factors (add):</p> <ul style="list-style-type: none"> Doxycycline 100mg PO BID Trimethoprim/sulfamethoxazole 5-8mg/kg/day for SSTI and 8-12mg/kg/day for osteomyelitis Linezolid 600mg PO BID Vancomycin (pharmacy consult) 	<p>7 – 14 days for soft tissue infection only</p> <p>If osteomyelitis:</p> <ul style="list-style-type: none"> 24-48 hours if complete resection of infected tissue 7-14 days if only residual soft tissue infection 21 days if residual bone infection with resection or 28-42 days without resection <p>Conversion to PO antibiotics (including osteomyelitis)[‡]:</p> <ul style="list-style-type: none"> Active agent available Functioning GI tract Clinically stable Select active agent(s) against recovered pathogen(s)

‡There is no prespecified duration of IV antibiotics required.

*Adjust antibiotic doses according to renal function as needed.

Classification	Pathogens	Empiric Antibiotic Therapy*	Duration of Treatment
4/Severe	<p>GPC (including MRSA) ± GNR ± anaerobes</p> <ul style="list-style-type: none"> • Cultures from the last 12 months may help empiric antibiotic selection. • Adjust empiric antibiotic regimen based on deep culture and susceptibility results 	<p>Preferred:</p> <ul style="list-style-type: none"> • Ceftriaxone 2g IV daily + metronidazole 500mg PO Q12H <p>PLUS</p> <ul style="list-style-type: none"> • Vancomycin (pharmacy consult), OR • Linezolid 600mg PO BID[‡] <p><i>P. aeruginosa</i> only with risk factors (change to):</p> <ul style="list-style-type: none"> • Piperacillin/tazobactam 4.5g IV Q8H • Cefepime 2g IV Q8H + metronidazole 500mg PO Q12H <p>[‡]Linezolid is preferred for use in necrotizing soft tissue infections</p>	<p>7 – 14 days for soft tissue infection only</p> <p>If osteomyelitis:</p> <ul style="list-style-type: none"> • 24-48 hours if complete resection of infected tissue • 7 – 14 days if only residual soft tissue infection • 21 days if residual bone infection with resection or 28 – 42 days without resection <p>Conversion to PO antibiotics (including osteomyelitis) [‡]:</p> <ul style="list-style-type: none"> • Active agent available • Functioning GI tract • Clinically stable • Select active agent(s) against recovered pathogen(s)

‡There is no prespecified duration of IV antibiotics required.

*Adjust antibiotic doses according to renal function as needed.

VI. References

1. Cortes-Penfield NW, Armstrong DG, Brennan MB, et al. Evaluation and management of diabetes-related foot infections. *Clin Infect Dis*. 2023;77(3):e1-e13. doi: 10.1093/cid/ciad255.
2. Gariani K, Lebowitz D, Kressmann B, et al. Oral amoxicillin-clavulanate for treating diabetic foot infections. *Diabetes Obes Metab*. 2019;21(6):1483-1486. doi: 10.1111/dom.13651.
3. Lipsky BA, Itani K, Norden C, and Linezolid Diabetic Foot Infections Study Group. Treating infections in diabetic patients: a randomized, multicenter, open-label trial of linezolid versus ampicillin-sulbactam/amoxicillin-clavulanate. *Clin Infect Dis*. 2004;38(1):17-24. doi: 10.1086/380449.
4. Lipsky BA and Uçkay I. Treating diabetic foot osteomyelitis: a practical state-of-the-art update. *Medicina (Kaunas)*. 2021;57(4):339. doi: 10.3390/medicina57040339.
5. McCreery RJ, Lyden E, Anderson M,, and Van Schooneveld TC. Impact of a syndrome-specific antibiotic stewardship intervention on antipseudomonal antibiotic use in inpatient diabetic foot infection management. *Antimicrob Steward Healthc Epidemiol*. 2023;3(1):e39. doi: 10.1017/ash.2023.123.
6. Mergenhagen KA, Croix M, Starr, KE, et al. Utility of methicillin-resistant *Staphylococcus aureus* nares screening for patients with a diabetic foot infection. *Antimicrob Agents Chemother*. 2020;64(4). doi: 10.1128/AAC.02213-19.
7. Morelli MK, Son AH, Bitar Y, and Hecker MT. Fear of missing organisms (FOMO): the discordance among broad-spectrum empiric antibiotic therapy, microbiologic results, and definitive antibiotic therapy for diabetic foot infections and lower extremity osteomyelitis. *Antimicrob Steward Healthc Epidemiol*. 2023;3(1):e186. doi: 10.1017/ash.2023.467.
8. Senneville É, Albalawi Z, van Asten SA, et al. IWGDF/IDSA guidelines on the diagnosis and treatment of diabetes-related foot infections (IWGDF/IDSA 2023). *Clin Infect Dis*. 2023. doi: 10.1093/cid/ciad527.
9. Pham TT, Gariani K, Richard JC, et al. Moderate to severe soft tissue diabetic foot infections: a randomized, controlled, pilot trial of post-debridement antibiotic treatment for 10 versus 20 days. *Ann Surg*. 2022;276(2):233-238. doi: 10.1097/SLA.0000000000005205.
10. Veve MP, Mercurio NJ, Sangiovanni RJ, et al. Prevalence and predictors of *Pseudomonas aeruginosa* among hospitalized patients with diabetic foot infections. *Open Forum Infect Dis*. 2022;9(7). doi: 10.1093/ofid/ofac297.

This document was reviewed and endorsed by the Vanderbilt Adult Antimicrobial Stewardship Committee on 7/17/2025


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Date last updated: 7/17/25
Date VASP approved: 7/17/25