

Evaluation of Local Pathogens and Management of Diabetic Foot Infections Jessica Foley (Seadler), PharmD,¹ Elizabeth Cady, PharmD, BCPS,² Praveen Mullangi, MD,³ Natalie Tucker, PharmD, BCPS, BCIDP⁴ 1. Dayton Children's Hospital, Dayton, OH 2. SIUE School of Pharmacy, Edwardsville, IL 3. Springfield Clinic LLC, Springfield, IL 4. HSHS St. John's Hospital, Springfield, IL

Background

- The prevalence of diabetes and its sequelae, including diabetic foot infection (DFI), are rising in the United States¹
- Complications of DFI such as amputation illustrate the importance of early and effective therapies such as antibiotics and surgery
- Per IDSA consensus guidelines, antimicrobial selection for DFI treatment should be guided by the severity of infection²
- □ This institution observed a lack of compliance with consensus guideline recommendations
 - Regardless of severity, hospitalized patients receive broad-spectrum antibiotics that included coverage for both methicillin-resistant *S. aureus* (MRSA) and *P. aeruginosa*

• CDC strongly recommends that antimicrobial stewardship programs create facility-specific guidance for common infections³

Study Purpose: Evaluate current practice with a goal of creating institutionspecific treatment guidance for providers to optimize the management of DFI

Objectives

Primary objective: Rate of guideline-compliant empiric antibiotic regimens

Secondary objectives:

Rate of empiric MRSA coverage 2. Rate of empiric *P. aeruginosa* coverage Duration of antibiotic therapy per patient

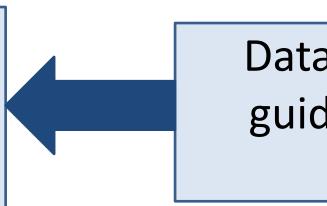
Methods

Patients identified by ICD-10 code for DFI between 8/1/18 – 7/31/19; n = 193

Exclusion criteria: 1. Antibiotics were continuation of outpatient therapy or being used for a concomitant infection 2. Any admission beyond the patient's first admission in study period 3. Pregnancy

114 patients included Data collection (retrospective chart review) Data compared with consensus guideline recommendations to determine compliance

Creation of institution-specific guidance for multidisciplinary management of DFI



References

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- 2. Lipsky BA, Berendt AR, Cornia PB, et al. 2012 Infectious Diseases Society of America Clinical Practice Guideline for the Diagnosis and Treatment of Diabetic Foot Infections. Clin Infect Dis 2012;54(12):132-73.
- 3. CDC. Core Elements of Hospital Antibiotic Stewardship Programs. Atlanta, GA: US Department of Health and Human Services, CDC; 2019. Available at <u>https://www.cdc.gov/antibiotic-use/core-elements/hospital.html</u>. 4. Lipsky BA, Aragon-Sanchez J, Diggle M, et al. IWGDF Guidance on the Diagnosis and Management of Foot Infections in
- Persons with Diabetes. Diabetes Metab Res Rev 2016;32(supplement 1):45-74.
- 5. Nelson A, Wright-Hughers A, Backhouse MR, et al. CODIFI (Concordance in Diabetic Foot Ulcer Infection): A Cross-Sectional Study of Wound Swab Versus Tissue Sampling in Infected Diabetic Foot Ulcers in England. BMJ Open 2018;8:e019437.

Methods (cont.)

Table 1: DFI Severity Classification, Expected Pathogens, and **Recommended Empiric Therapy²**

IDSA Infection Severity	Pathogens	Antibiotics Recommended	
Mild (local infection	MSSA,	Dicloxacillin, clindamycin,	
involving only the	Streptococcus spp.	cephalexin, amox-clav	
skin/subcutaneous tissue)	MRSA	Doxycycline, SMX-TMP	
Moderate (involvement of deeper structures – abscess, osteomyelitis, septic arthritis, fasciitis – without	MSSA, <i>Streptococcus</i> spp., Enterobacteriaceae, anaerobes	Levofloxacin, ceftriaxone, amp-sulbactam, moxifloxacin, ertapenem, ciprofloxacin w/ clindamycin	
signs of systemic response)	MRSA	Linezolid, vancomycin, daptomycin	
Severe (moderate infection and meeting > 2 SIRS criteria)	P. aeruginosa*	Pip-tazo, cefepime, ceftazidime, aztreonam, meropenem	

*coverage indicated in severe infections or in patients with validated risk factors, such as previous isolation of *P. aeruginosa* or frequent foot soaking

Table 2: Baseline Characteristics

Characteristic

Age, years, mean (SD)

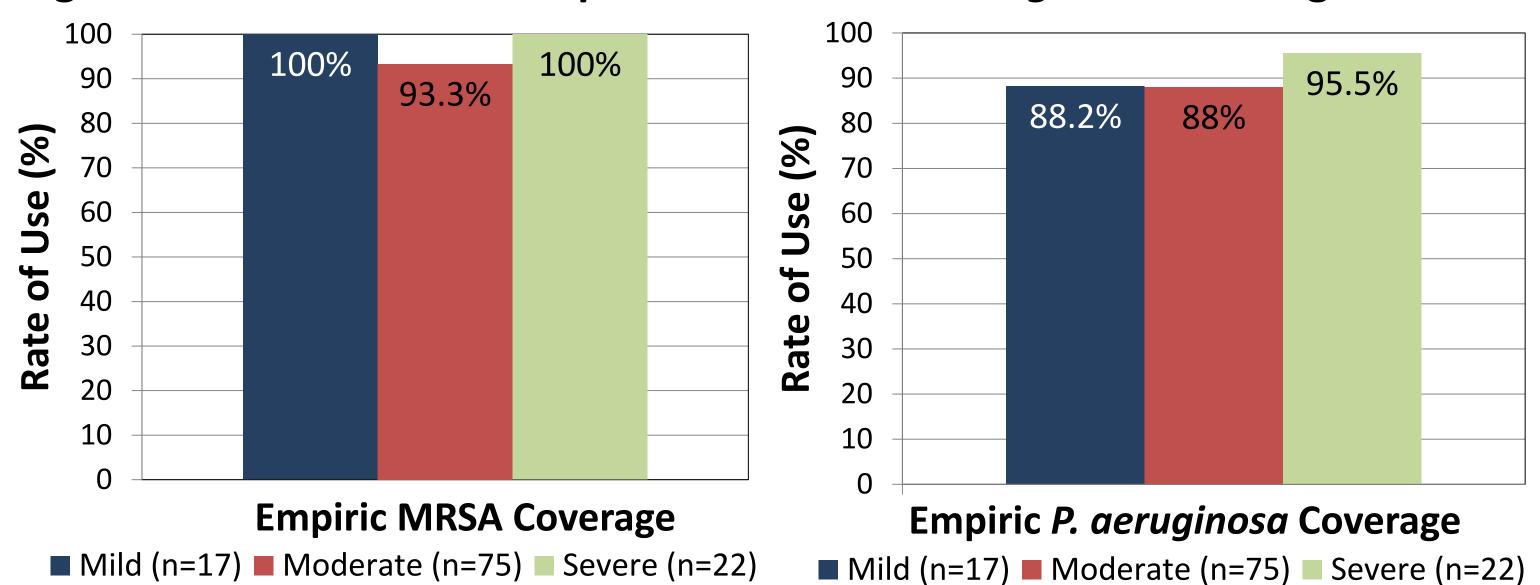
Male, n (%)

Hemoglobin A1c% during index encour

Table 3: Rate of Guideline-Compliant Empiric Antibiotic Regimens

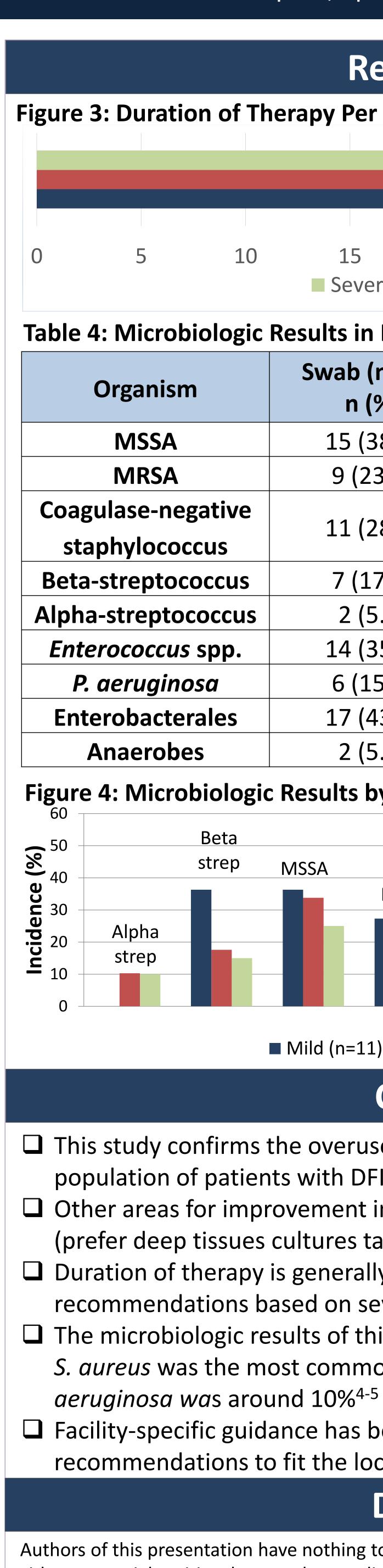
	IDSA Infection Severity			
Guideline- Compliant Empiric Regimen	Mild	Moderate	Severe	$T_{otol} (n - 114)$
	(n=17),	(n=75) <i>,</i>	(n=22),	Total (n=114),
	n (%)	n (%)	n (%)	n (%)
Yes	0 (0)	8 (10.67)	22 (100)	30 (26.3)
Νο	17 (100	67 (89.3)	0 (0)	84 (73.7)
		p = 0.0001		

Figures 1 and 2: Rates of Empiric MRSA and *P. aeruginosa* Coverage



Results

	Ν
	62 (<u>+</u> 10.8)
	81 (71)
inter, mean (SD)	9 (<u>+</u> 2.2)

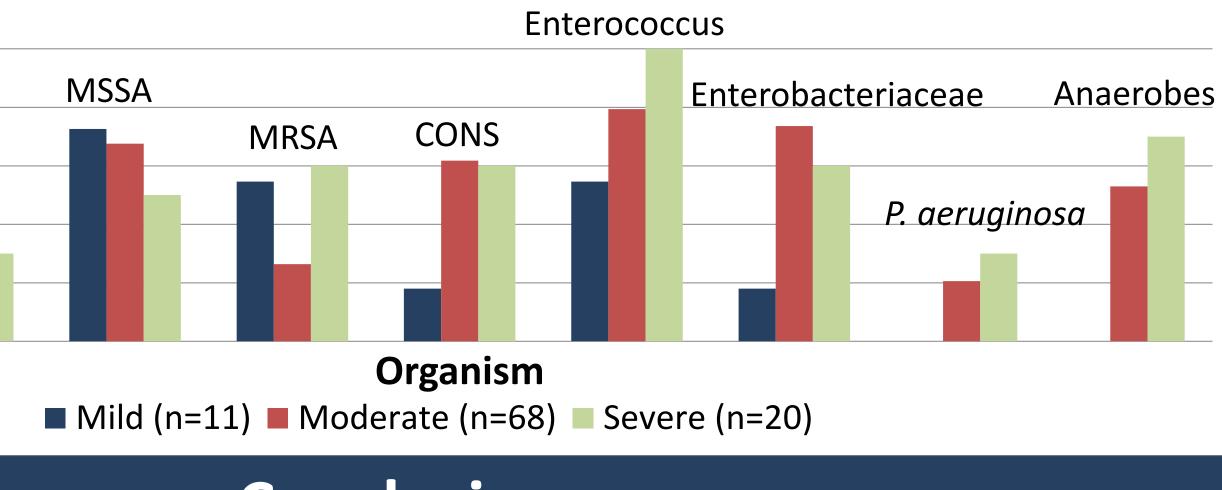


Results (cont.)						
f Therapy Per Patient (Days)						
				29.7		
		21.3			35.2	
10	15 Severe	20 Moderate	25 ■ Mild	30	35	40

Table 4: Microbiologic Results in Full Population

	Swab (n=39), n (%)	Deep tissue or surgical (n=60), n (%)	All cultures (n=99), n (%)
	15 (38.5)	17 (28.3)	32 (32.3)
	9 (23.1)	9 (15)	18 (18.2)
e	11 (28.2)	17 (28.3)	28 (28.3)
S	7 (17.9)	12 (20)	19 (19.2)
JS	2 (5.1)	7 (11.7)	9 (9.1)
•	14 (35.8)	26 (43.3)	40 (40.4)
	6 (15.4)	4 (6.7)	10 (10.1)
	17 (43.6)	15 (25)	32 (32.3)
	2 (5.1)	23 (38.3)	25 (25.3)

Figure 4: Microbiologic Results by Severity of Infection



Conclusion

This study confirms the overuse of broad-spectrum antibiotics in this

- population of patients with DFI, particularly *P. aeruginosa* coverage
- Other areas for improvement include culture collection method and timing
 - (prefer deep tissues cultures taken prior to antibiotic initiation)
- Duration of therapy is generally consistent with consensus guideline recommendations based on severity of infection
- □ The microbiologic results of this study are concordant with previous studies, as S. aureus was the most common organism isolated and the rate of P.
- □ Facility-specific guidance has been created by focusing consensus guideline recommendations to fit the local need based on the results of this study

Disclosures

Authors of this presentation have nothing to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation.

Please send any correspondence to foleyj1@childrensdayton.org.