Antibiotic Susceptibility Patterns of Commonly Isolated Bacteria July 2024 – June 2025 (12 months)

All MMC Sites

NOTES

- 1. Minimum inhibitory concentrations (MIC) and interpretations are based on the CLSI standards and an advanced antibiotic expert system.
- 2. Percentages are not calculated for organisms with <10 isolates. For *N* of < 30 isolates, results may not be statistically relevant. Interpret with caution.

Box color: Intrinsic Resistance Less susceptible More susceptible

Text color: • > 10% increase in susceptibility from previous year • > 10% decline in susceptibility from previous year

	Αľ	MPI	CEF	TRIAX	CIPR	OFLX	TMP	/SMX
	N	% S	N	% S	N	% S	Ν	% S
Salmonella species (all inpatient isolates) ²	29	97	18	94	29	62	29	100

	Р	EN	CEF	TRIAX	VANC		
	N	% S	N	% S	N	% S	
viridans Streptococcus (sterile sites)	66	77	68	99	68	100	

STREPTOCOCCUS		Ster	ile Site		Non-Sterile Site						
All Campuses	2024-2025	N	S	_	R	N	S	1	R		
	Meningitis	44	68		32						
A B	Non-CNS	44	91	9	0						
PENICILLIN ^{A,B}	Parenteral					75	84	8	8		
	Oral					75	49	19	32		
ossetnia von s ^A	Meningitis	44	93	5	2	75	91	1	8		
CEFTRIAXONE ^A	Non-CNS	44	98	2	0	75	92	8	0		
LEVOFLO)	KACIN	47	98	0	2	77	99	0 1			
TRIMETH/S	SULFA ^C					76	66	11	24		

A. Pneumococcal susceptibility rates against penicillin and ceftriaxone from sterile sites are reported as if isolates came from both CSF and all other sterile sites. Susceptibility rates are higher for non-CSF sites because higher antibiotic concentrations can be reached.

B. For pneumococcal isolates from non-sterile sites (sputum), penicillin susceptibility rates are also reported separately for oral and parenteral formulations. The suceptibility rate is higher for parenteral than oral penicillin because higher concentrations are achieved when penicillin is given parenterally.

C. Pneumococci from sterile sites are not tested against erythromycin and trimethoprim-sulfamethoxazole because those antimicrobials generally should be used only for pneumococcal respiratory infections.

ENTEROCOCCUS Sterile Sites	A	AMPI	D	APTO ^A	GEN	NT SYN ^B	L	INEZD	STR	EP SYN ^B	V	/ANC
All Campuses 2024-2025	Ν	% S	Ν	% S	N	% S	N	% S	Ν	% S	Ν	% S
Enterococcus faecalis	139	99	139	67	139	77	139	99	139	87	139	92
Enterococcus faecium	96	10	94	95	95	86	96	98	95	60	94	36

A. For *E. faecalis*, daptomycin is not recommended due to cost and the availability of an agent with a narrower spectrum of activity (i.e. ampicillin/amoxicillin).

B. Susceptibility indicates synergy with penicillin, ampicillin, piperacillin-tazobactam, and vancomycin.

CANDIDA All Campuses	C. albicans					C. parapsilosis ²					C. tropicalis ²					C. glabrata					C. auris ^{A,2}		
2024-2025	N	S	SDD	_	R	Ν	S	SDD	ı	R	Ν	S	SDD	-	R	Ν	S	SDD	_	R	N	S	R
Fluconazole	87	95	2		2	18	89	11		0	22	41	45		14	41		93		7	19	11	89
Voriconazole	87	98		1	1	18	100		0	0	22	41		55	5								
Micafungin	49	98		0	2	12	17		0	83	18	94		0	6	39	100		0	0	19	100	0
Amphotericin B																				19	47	53	

*Data is shown for epidemiologic purposes; contact ID for questions about use of antifungals.

A. Breakpoints for *C. auris* have not been established by CLSI. Breakpoints used here are defined by the CDC and are based on those established for closely related *Candida* species and on expert opinion.