Antibiotic Susceptibility Patterns of Commonly Isolated Bacteria July 2023 – June 2024 (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	27	93	7	2	26	73	27	96	

	Р	EN	CEF	TRIAX	VANC		
	N	% S	N	% S	N	% S	
viridans Streptococcus (sterile sites)	57	70	62	97	60	98	

STREPTOCOCCUS I	PNEUMONIAE		Steri	ile Site			Non-St	erile Site			
All Campuses	N	S	_	R	N	S	ı	R			
	Meningitis	52	62		38						
DENIGH LINA.B	PENICILLIN ^{A,B} Non-CNS 52 98 0 Parenteral	2									
PENICILLIN'	Parenteral					65	91	6	3		
	Oral					65	62	15	23		
CEETDIA VON EA	Meningitis	53	98	2	0	65	95	2	3		
CEFTRIAXONE ^A	Non-CNS	53	100	0	0	65	97	15 23	3		
LEVOFLO)	(ACIN	60	96	2	2	72	94 3				
TRIMETH/S	SULFA ^C					71	76	76 8 1			

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	IT SYN ^B	LI	NEZD	STR	EP SYN ^B	VANC		
All Campuses 2023-2024	N	% S	N	% S	N	% S	N	% S	N	% S	N	% S	
Enterococcus faecalis	143	100	143	85	143	76	143	99	143	85	143	93	
Enterococcus faecium	93	11	92	96	92	89	93	98	92	59	93	31	

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}		
2023-2024	N	S	SDD	1	R	N	S	SDD	ı	R	Ν	S	SDD	ı	R	N	S	SDD	ı	R	Ν	S	R
Fluconazole	88	93	2		3	23	87	9		13	12	42	33		25	56		82		18	21	5	95
Voriconazole	88	93		3	3	23	91		9	0	12	42		58	0								
Micafungin	5	2		2	2	6	2		2	2	3	2		2	2	56	96		0	4	20	100	0
Amphotericin B																					20	90	10

*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.