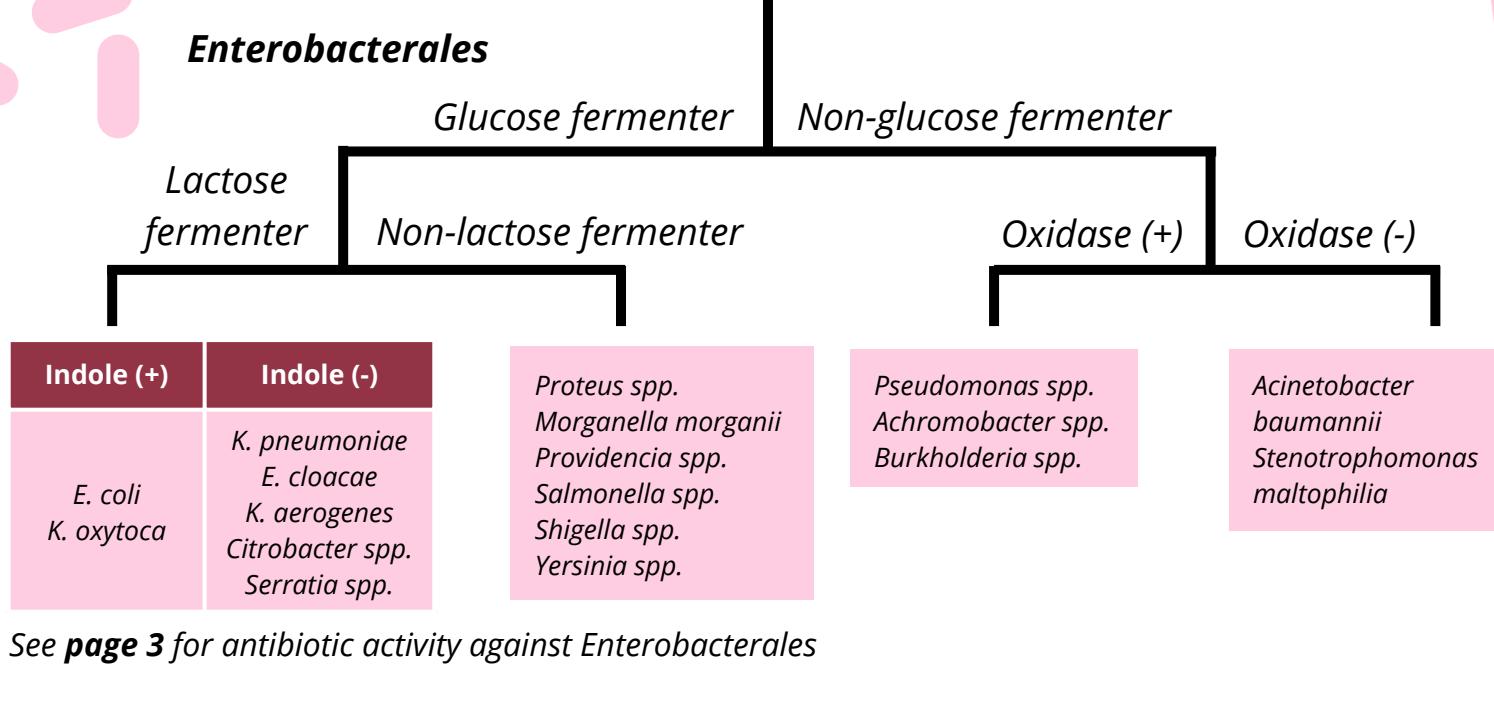


Gram-Negative Bacilli

Disclaimer: This document is for educational purposes and not to guide patient-level therapeutic decisions



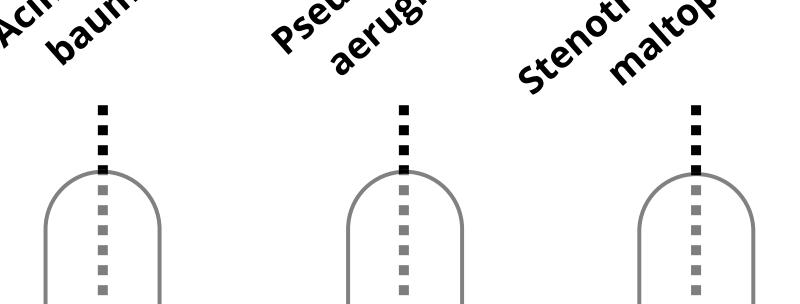
See **page 3** for antibiotic activity against **Enterobacteriales**

KEY

- Routinely active
- Variable activity
- No or limited activity

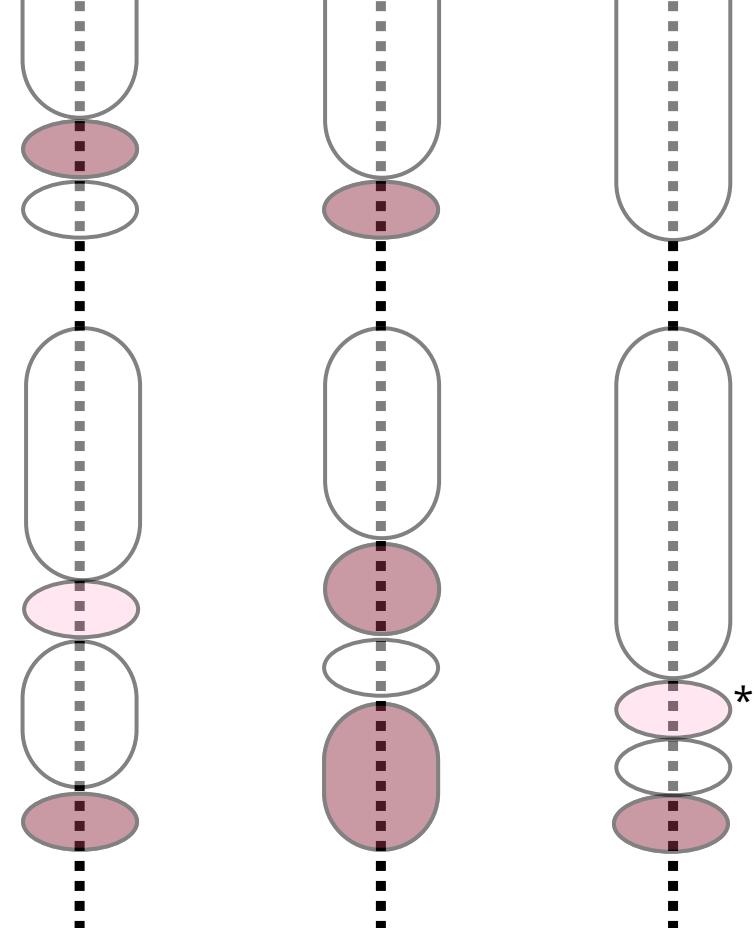
Penicillins

- Penicillin G
- Anti-staphylococcal penicillins (OXA/NAF)
- Ampicillin/Amoxicillin
- Amoxicillin-clavulanate
- Ampicillin-sulbactam
- Piperacillin-tazobactam



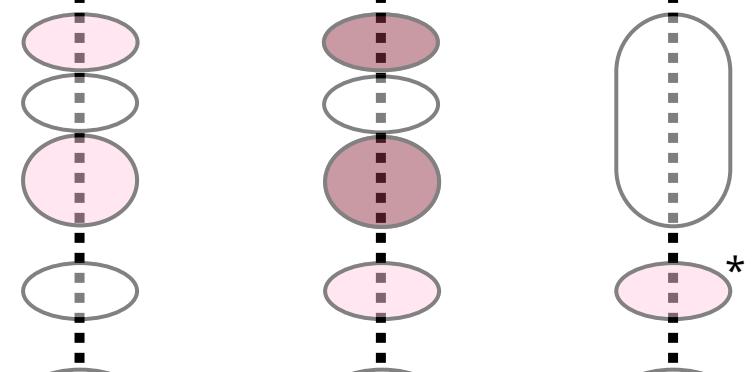
Cephalosporins

- 1st Gen (LEX/CFZ)
- 2nd Gen (Cefuroxime)
- 3rd Gen
- Ceftriaxone/Cefpodoxime
- Ceftazidime
- 4th Gen (Cefepime)
- Ceftaroline
- Ceftazidime/avibactam*
- Ceftolozane/tazobactam
- Cefiderocol

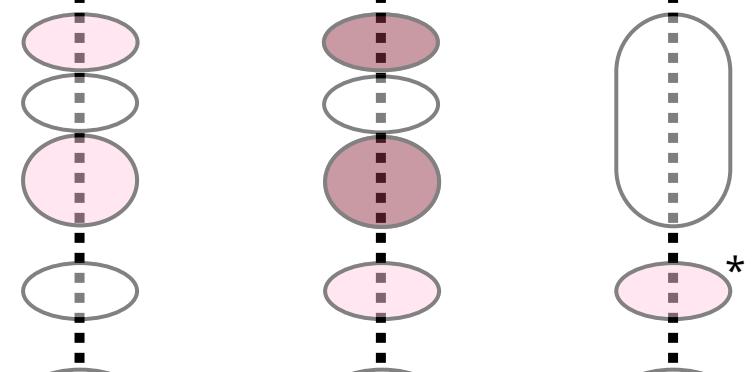


Carbapenems

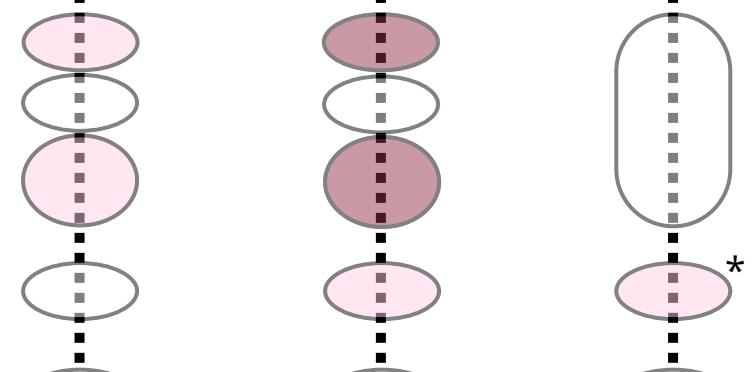
- Meropenem/Imipenem
- Ertapenem
- Meropenem/vaborbactam
- Imipenem/relebactam



Aztreonam*

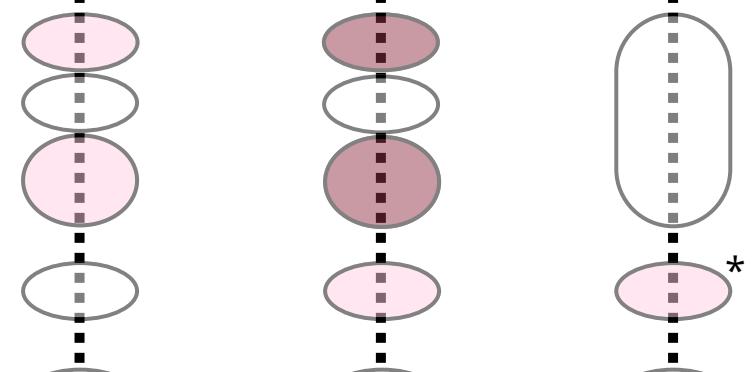


Macrolides



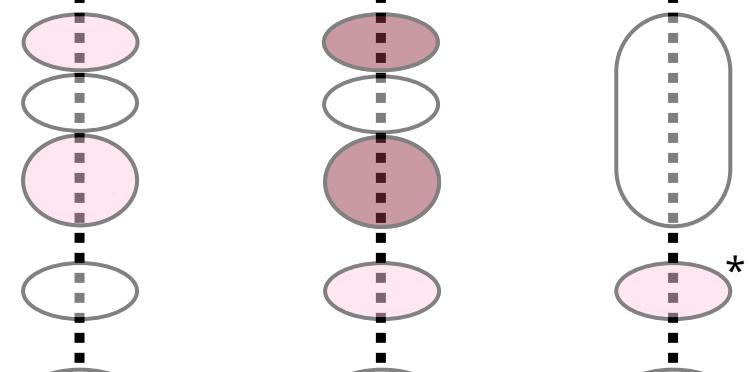
Tetracyclines**

- Doxycycline
- Minocycline
- Ervacacycline/
- Tigecycline

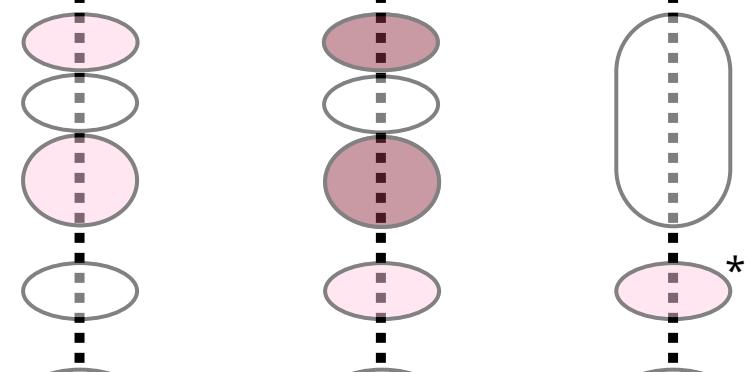


Fluoroquinolones

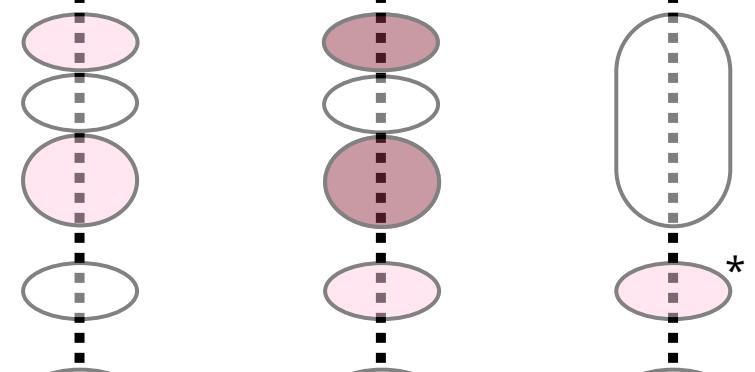
- Levofloxacin
- Ciprofloxacin
- Moxifloxacin



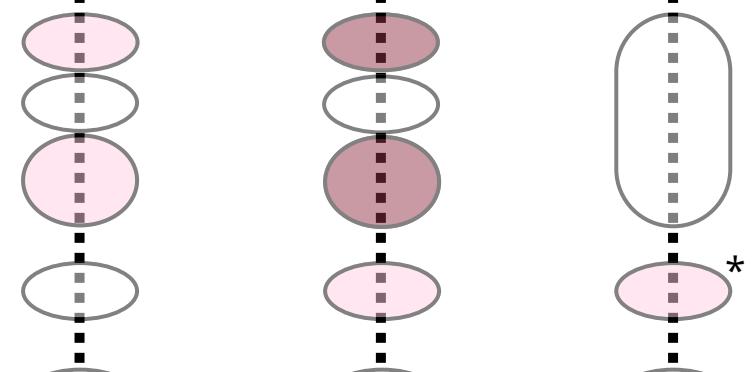
Aminoglycosides



TMP/SMX



Colistin/Polymyxin B***



*Ceftazidime/avibactam should be given in combination with aztreonam for moderate to severe *S. maltophilia* infections; **Minocycline is the preferred tetracycline for *A. baumannii* infections. Tigecycline may be considered as an alternative. Ervacacycline demonstrates in vitro activity but there is limited clinical data to support its use;

***Given significant toxicity and availability of alternative agents, colisitin and polymyxin B should be considered as a last-line treatment option in most clinical scenarios

Enterobacterales Resistance -- An Overview of Common Terminology

	"AmpC"	"ESBL"	"CRE"
Description	Inducible expression of the ampC beta-lactamase among certain Enterobacterales organisms (AmpC-E)	Presence of extended-spectrum beta-lactamase (ESBL) gene expression	Carbapenem-resistant Enterobacterales (CRE) includes resistance via carbapenemase-producing genes (see below) or other resistance mechanisms
Genotype	Many but not commonly tested in the clinical setting	CTX-M , SHV, TEM	KPC , NDM/IMP/VIM (metallo-beta-lactamases), OXA-48
Phenotype	3rd generation cephalosporin-resistant OR resistance develops with antibiotic exposure	Ceftriaxone-resistant	Resistance to ≥ 1 carbapenem (<i>i.e.</i> , may be ertapenem-resistant but meropenem-susceptible)
Typical Organisms	HECK-Yes <i>(Hafnia alvei, Enterobacter cloacae, Citrobacter freundii, Klebsiella aerogenes, Yersinia enterocolitica)</i>	<i>E. coli, K. pneumoniae, K. oxytoca, P. mirabilis</i>	KPC -- <i>K. pneumoniae</i> Non-carbapenemase-producing, carbapenem-resistant Enterobacterales -- Enterobacter spp., K. pneumoniae, E. coli

KEY

- Routinely active
- Variable activity
- No or limited activity

Enterobacterales
(no ESBL, ampC, or carbapenemase production)

Penicillins

Penicillin G
Anti-staphylococcal penicillins (OXA/NAF)
Ampicillin/Aoxicillin
Amoxicillin-clavulanate
Ampicillin-sulbactam
Piperacillin-tazobactam

ampC-Enterobacterales
ESBL-producing Enterobacterales

Cephalosporins

1st Gen (LEX/CFZ)
2nd Gen (Cefuroxime)
3rd Gen
 Ceftriaxone/Cefpodoxime
 Ceftazidime
4th Gen (Cefepime)
Ceftaroline
Ceftazidime/avibactam*
Ceftolozane/tazobactam
Cefiderocol

CRE (KPC)

CRE (NDM/IMP/VIM)

CRE (OXA-48)

Carbapenems

Meropenem/Imipenem
Ertapenem
Meropenem/vaborbactam
Imipenem/relebactam

CRE (NDM/IMP/VIM)

CRE (OXA-48)

Aztreonam*

CRE (OXA-48)

Macrolides

CRE (OXA-48)

Tetracyclines

Doxycycline
Minocycline
Eravacycline/
Tigecycline

CRE (OXA-48)

CRE (OXA-48)

Fluoroquinolones

Levofloxacin
Ciprofloxacin
Moxifloxacin

CRE (OXA-48)

CRE (OXA-48)

Aminoglycosides

CRE (OXA-48)

TMP/SMX

CRE (OXA-48)

*Ceftazidime/avibactam should be given in combination with aztreonam for CRE due to metallo-beta-lactamases; **Given significant toxicity and availability of alternative agents, colisitin and polymyxin B should be considered as a last-line treatment option in most clinical scenarios