



## Beat the Bug: *Pseudomonas aeruginosa*

*Pseudomonas aeruginosa* is non-fermenting gram-negative bacilli that is associated with healthcare exposure. Preserving *P. aeruginosa* antibiotic activity through de-escalation of spectrum or discontinuation of unnecessary antibiotics is a key antimicrobial stewardship activity. *P. aeruginosa* is intrinsically resistant to many antibiotics and acquired resistance is commonly encountered. Which antibiotics have activity against *P. aeruginosa* and what are some considerations for use?

	Antibiotic	Notes
<b>First Line</b>	Piperacillin-tazobactam	-Pip-tazo has activity against <a href="#">E. faecalis</a> and <a href="#">Bacteroides spp.</a> whereas cefepime and ceftazidime do not - <a href="#">Lower risk of C. difficile</a> infection than other anti-pseudomonals
	Ceftazidime, Cefepime	-Cefepime stable against <a href="#">AmpC</a> - <a href="#">Ceftazidime less reliable against S. pneumoniae and MSSA</a> - <a href="#">Both can be used in patients with penicillin allergies, including anaphylaxis</a>
	<a href="#">Aztreonam</a>	-Limit to patients with allergies to alternatives
	Meropenem, Imipenem	- <a href="#">Ertapenem is NOT active</a> -Non-carbapenem beta-lactams preferred, if active, to preserve carbapenem activity for infections due to other resistant bacteria (e.g. <a href="#">ESBLs</a> )
	Ciprofloxacin, Levofloxacin, Delafloxacin	-Quinolones are <a href="#">ONLY oral options</a> -Moxifloxacin is NOT active against <i>P. aeruginosa</i>
	Tobramycin, Amikacin, Plazomicin	-Gentamicin breakpoints recently retired, so no longer considered active -Avoid as monotherapy <a href="#">except in UTIs</a> due to inferiority
<b>If Resistant to Above</b>	Ceftolozane-tazobactam Ceftazidime-avibactam Imipenem-relebactam	-Ceftolozane-tazobactam preferred, if active, to preserve caz-avi and imi-rel for infections due to other resistant bacteria (e.g. <a href="#">KPCs</a> )
	Cefiderocol	-Novel beta-lactam/beta-lactamase inhibitors preferred, if active, to preserve cefiderocol activity against other resistant bacteria (e.g. <a href="#">metallo-β-lactamase</a> producing organisms)
<b>Last Line</b>	Polymyxin B, Colistin <sup>2</sup>	-No susceptible category, highlighting role only as last line -Polymyxin B preferred over colistin in systemic infections due to better kinetics and lower nephrotoxicity risk -Do not use polymyxin B in UTIs due to no urinary distribution
<b>Do Not Use</b>	Cefepime-enmetazobactam Meropenem-vaborbactam	Enmetazobactam and vaborbactam do NOT restore meaningful activity in <i>P. aeruginosa</i> that are resistant to cefepime and meropenem <sup>3,4</sup>

**Key Takeaway:** Antibiotics with activity against *P. aeruginosa* are limited and often are last line options for other resistant bacteria. Reserving anti-pseudomonal antibiotics whenever possible is an important antimicrobial stewardship target.

### References

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