

Drug	Spectrum	MOA	Pharmacokinetics	ADRs
1-Penicillins Note: first 3 drugs (3 rows) are <i>Inhibitors of cell wall synthesis</i> (β-Lactam Antibiotics)	<i>Extended Spectrum Penicillins:</i> Amoxicillin, Ampicillin -Active against gram positive & gram negative microorganism. <i>Narrow Spectrum Penicillin</i> <i>Penicillin G:</i> -Narrow spectrum -Destroyed by gastric acidity (so it is given parentally) -Inactivated by β - lactamase -Short acting (4-6 hrs)	-Irreversibly inhibits transpeptidase enzyme that catalyze the final step in cell wall synthesis of bacteria. (Inhibits the synthesis of peptidoglycan layer of bacterial cell wall)	-Inactivated by β- lactamase enzyme, (So given with β-lactamase inhibitors are available <u>e.g Amoxicillin + Clavulanic acid and ampicillin + salbactam.</u> -Amoxicillin and ampicillin are Acid stable(effective orally) -Can be given parenterally (I.V or I.M) -Amoxicillin is better absorbed from the gut & not affected by food.	-Hypersensitivity -Diarrhea -Nephritis -Neurotoxicity
<i>2-Cephalosporins</i> 3rd generation Ceftazidime Ceftriaxone	-Highly effective against Gm -ve bacilli -Anaerobic microbes -Pseudomonas -Highly resistant to β- lactamase compared to penicillin -Effective in Gm-ve meningitis	-Inhibits bacterial cell wall synthesis	-Both of them are given by intravenous infusion	-Allergy -Thrombophlebitis -Renal toxicity -Superinfections
<i>3-Carbapenems</i> Imipenem	-Has a wide spectrum of activity -Resistant to <u>most β lactamases</u> except metallo-β lactamase . 	-Bactericidal, inhibit bacterial cell wall synthesis.	-Not absorbed orally, taken by I.V. -Inactivated by dehydropeptidases in renal tubules, so it is given with an inhibitor cilastatin for clinical use. -Penetrates body tissues and fluids including C.S.F.	-Nausea, vomiting, diarrhea -Skin rash and reaction at the site of infusion -High doses in patients with renal failure may lead to seizures -Patients allergic to penicillins may be allergic to carbapenems .

<p>4-Vancomycin</p> <p>-May be combined with ampicillin or ceftazidime as an initial therapy of meningitis in infant, elderly and immunocompromised patients .</p>	<p>-Active only against Gm+ve bacteria</p> <p>-Used in combination with 3rd generation cephalosporins for treatment of meningitis caused by penicillin resistant pneumococci.</p> <p>-Good drugs used against(MRSA).</p>	<p>-Cell wall inhibitor</p>	<p>-Poorly absorbed orally</p> <p>-Given intravenously</p> <p>-Used orally to treat GIT infections caused by clostridium defficile e.g colitis.</p>	<p>-Phlebitis</p> <p>-Ototoxicity</p> <p>-Nephrotoxicity</p> <p>-Histamine release [red man (red neck)]</p> <p>-Hypotension</p>
<p>Fluoroquinolones</p> <p>Ciprofloxacin</p> <p>Contraindicated in :</p> <ul style="list-style-type: none"> ▶ Growing children (below 18 years) ▶ Pregnancy ▶ Lactation ▶ History of epilepsy or CNS disorder 	<p>-Effective against :Gm-ve organisms</p> <p>-Limited activity :against Gm+ve organisms</p> <p>-Effective in patients who are allergic to penicillins.</p> <p>-Effective against :intracellular pathogens such as: Legionella, Chlamydia, some mycobacteriae</p>	<p>Block bacterial DNA synthesis by inhibiting bacterial topoisomerase 11(DNA gyrase) & topoisomerase1V</p>	<p>-Well absorbed orally</p> <p>-Absorption is impaired by divalent cations ; iron, zinc or those in antacids as aluminium, magnesium)</p> <p>-Half-life 3hrs</p> <p>-Widely distributed in body fluids & tissues</p> <p>-Penetrates into CSF</p> <p>-Highly concentrated in bone, kidney, prostate, lung</p> <p>-Excreted through kidney & appear in breast milk</p>	<p>-GIT upset</p> <p>-CNS :Headache , dizziness, insomnia</p> <p>-Abnormal liver function tests</p> <p>-Skin rash & photosensetivity</p> <p>-Cartilage damage (arthropathy)</p> <p>-Tendon damage (tendinitis)</p> <p>-Enzyme inhibitor</p>