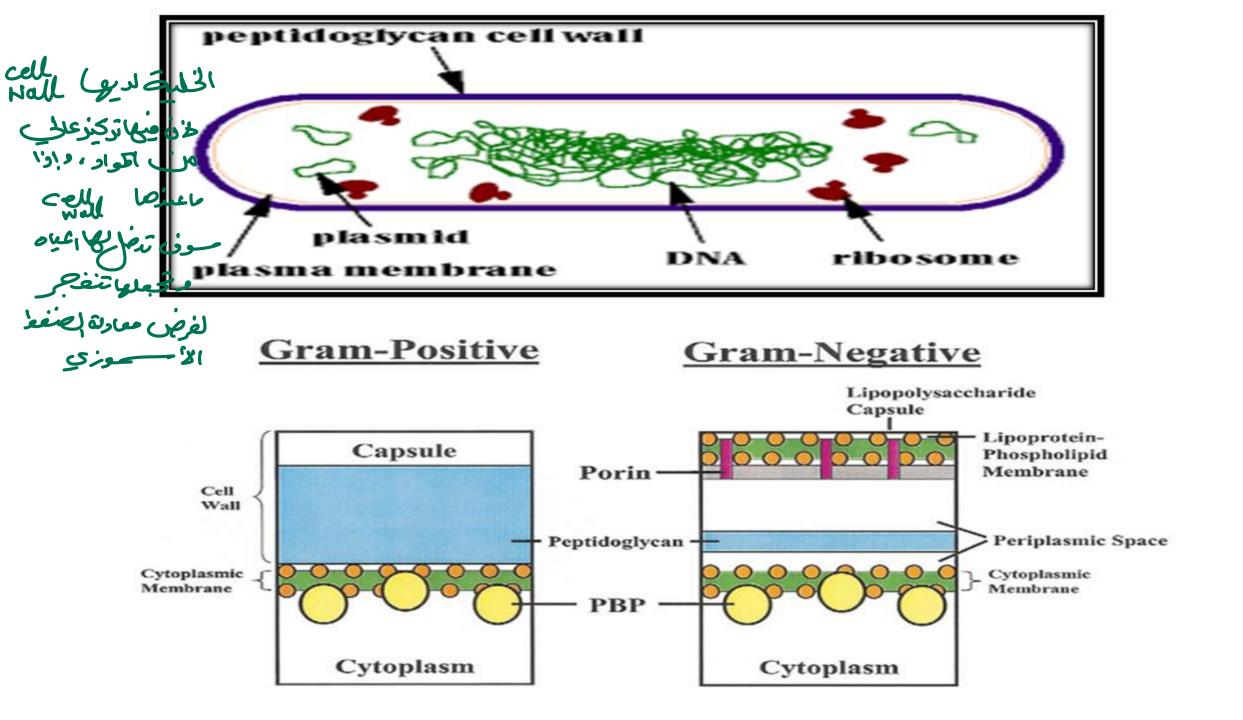
بسم الله الرحمن الرحيم

Pharmacology of antibacterial drugs Cell wall inhibitors (part 1)

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The major cell wall synthesis inhibitors currently in use are:

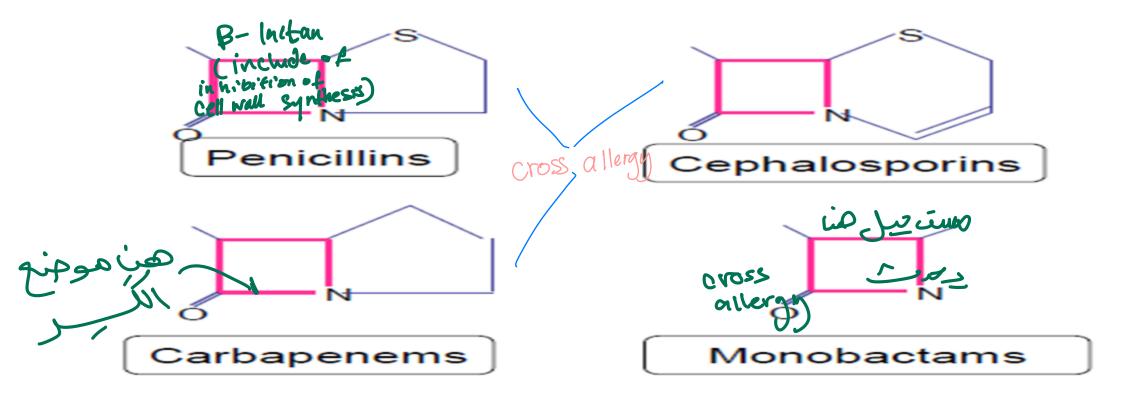
1- The **beta-lactams** (e.g., penicillin and cephalosporins), which block the formation of the peptidoglycan layer.

2- The **glycopeptides** (vancomycin and teicoplanin), which disrupt assembly of the peptidoglycan precursor lipid II.

Cell wall biosynthetic stages	Antibiotics	Target
Stage I: the cytoplasmic stage	D-Cycloserine	D-Ala–D-Ala ligase, alanine racemase
•	Fosfomycin	MurA
Stage II: the membrane-associated stage	Uridyl peptides (tunicamycin)	MraY
	Ramoplanin	MurG, lipid II
Stage III: the extracytoplasmic stage	β-Lactams	PBPs
	Glycopeptides	Lipid II (D-Ala–D-Ala terminal)
	Moenomycin	Transglycosylase
	Mannopeptimycins	Lipid II
	Lantibiotics (nisin)	Lipid II
	Defensin <mark>(</mark> plectasin)	Lipid II
	Bacitracin	Undecaisoprenyl pyrophosphate

Beta lactam antibacterial drugs

The β -lactams include **penicillins**, **cephalosporins**, **monobactams** and **carbapenems** and they share a common structure, and a common mechanism of action.



Features of beta lactam antibiotics:

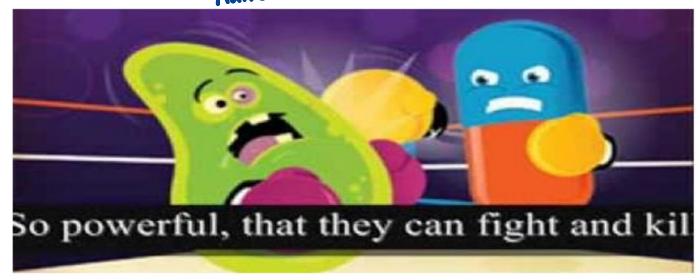
They contain the 4-membered ring (lactam) which is intrinsically labile to hydrolysis (acidic or enzymatic).

> Target: cell-wall biosynthesis > Complete natural resistance for all B-lactons

>Action: bactericidal, active only against growing cells.

➤They have variable spectrum.

Broach Tklm narrow





Acl side

chain

Beta-Lactam

Ring

General Structure of Penicillins

(Gluiaa'd)

Thiazolidine

Ring

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Chemistry:

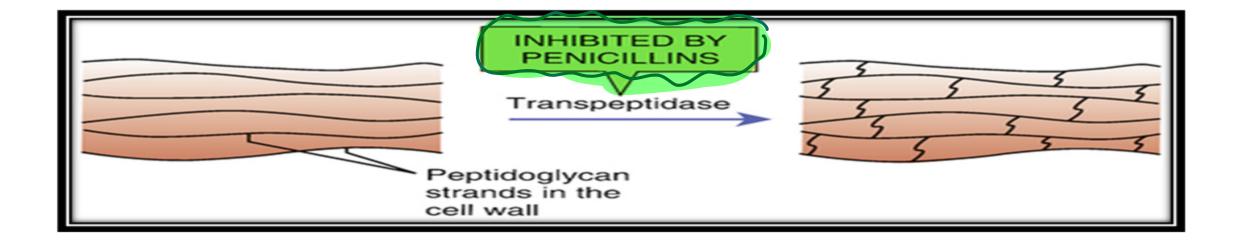
> The basic structure of the penicillins consists of a **thiazolidine ring** (A) connected to a β -lactam ring (B) to which is attached a side chain (R). > The β -lactam ring is responsible for the biological activity of penicillins, and it is targeted by organisms that produce penicillinase enzyme to destroy it.

mitated

➤The side chain (R) can be cleaved by amidase enzyme producing 6-aminopenicillanic acid to which new side chains can be added to produce new compounds of semi-synthetic penicillins. **Mechanism of action**

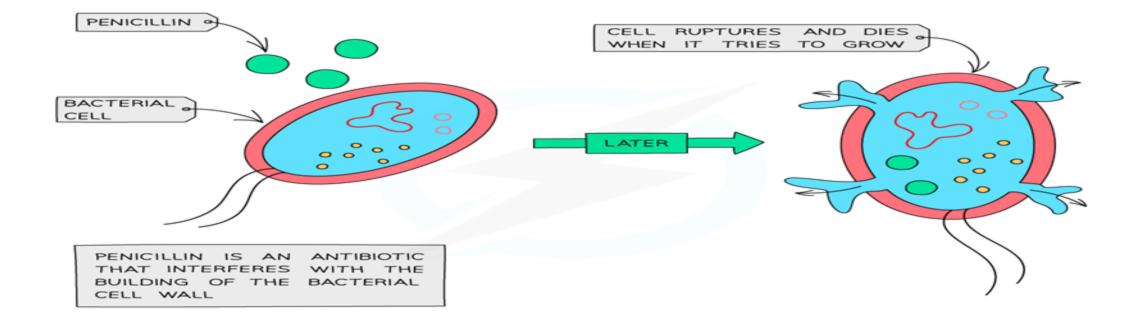
➢Penicillins are bactericidal through <u>inhibition of bacterial cell</u> wall synthesis for growing bacteria.

The bacterial cell wall consists of glycopeptides linked via five peptide bridges between amino acid side chains.
 Bacterial cells with evident cell wall have penicillin binding proteins (PBP) to which transpeptidases are attached (in the periplasmic space).



➤This trans-peptidation reaction gives the rigid mechanical stability of the cell wall and prevent osmotic shock.

➢Binding of Penicillins and other beta lactam drugs to PBP causes inhibition of these transpeptidases and inhibition cell wall synthesis occur leading to bacterial cell death.



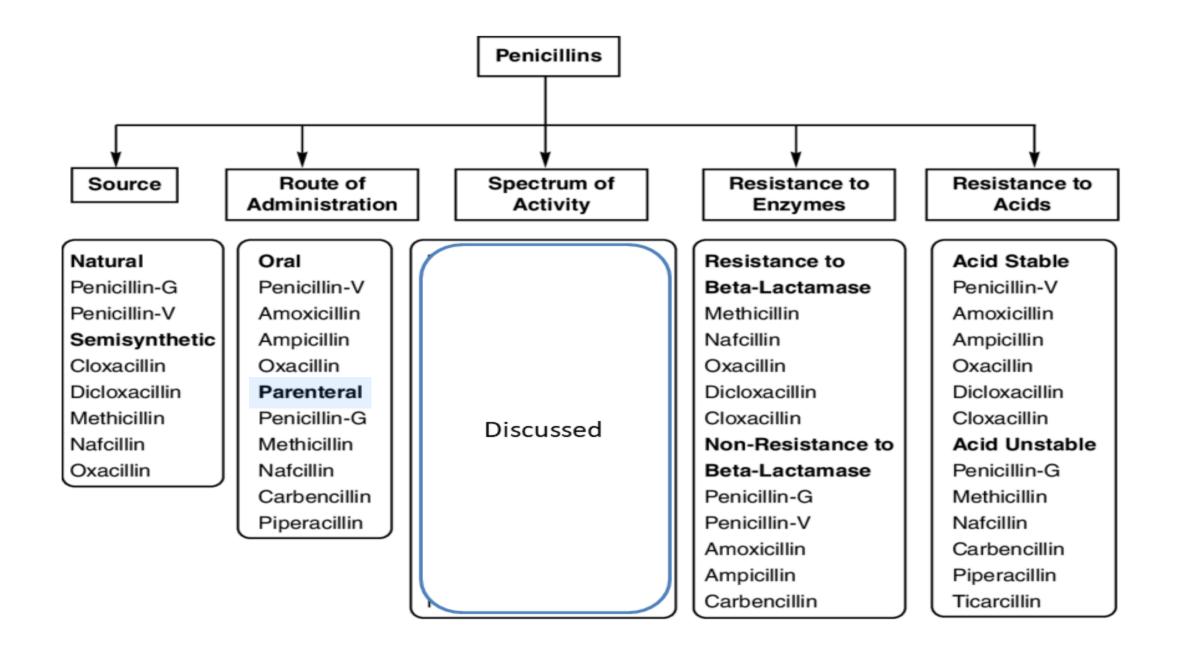
Mechanism of resistance to penicillins

- Enzymatic hydrolysis where bacteria produce β-lactamases
 (penicillinases) enzymes that can destroy β-lactam antibiotics.
- 2. Inability of the drug to penetrate to its site of action especially in gram negative bacteria.
- 3. Active efflux pumps that remove the antibiotic from its site of action.
- 4. Alteration in PBP with decreased affinity for β -lactam antibiotics.
- Natural (intrinsic) resistance. in bacteria lacking cell wall like Mycoplasma..

Classification of the penicillins

According to spectrum

The PENICILLINS Narrow spectrum penicillins Penicillin G Penicillin V Broad Spectrum Penicillins (aminopenicillin) Amoxicillin Ampicillin – Bacampicillin Penicillinase-resistant Penicillin (anti-staphyloccocal penicillins) Cloxacillin Nafcillin Methicillin Dicloxacillin Oxacillin Extended-Spectrum penicillins (Anti-pseudomonal penicillins) Carbenicillin Mezlocillin Piperacillin Ticacillin



<u>1- Narrow spectrum (natural) penicillins</u>

e.g. Natural Penicillins including penicillin (benzyl penicillin)
& penicillin (phenoxymethyl penicillin):
> Highly active against sensitive strains of gram-positive cocci, but they are readily hydrolyzed by penicillinase.
> May low spectrum
> They are ineffective against most strains of Staph. aureus.
> Some gram-negative cocci and anaerobic bacteria are susceptible to natural penicillins.

is rout of adminstration

2- The penicillinase resistant penicillins (Anti-staph penicillins)
 e.g. Methicillin, Nafeillin, Oxacillin, Cloxacillin, and Dicloxacillin.
 > They have very narrow spectrum (only active against sensitive strains of staphylococci), so, they are the agents of first choice for treatment of penicillinase-producing Staph aureus and Staph epidermidis that are not Methicillin resistant.

≻They are ineffective against bacilli and gram-negative organisms.

3- Broad spectrum penicillins (Aminopenicillins) e.g. **ampicillin** and **amoxicillin** which antimicrobial activity covers not only grampositive cocci but also the gram-negative organisms like Hemophilus influenza, E coli and proteus mirabilis. These drugs are administered frequently, with a β -lactamase inhibitor such as clavulanate or sulbactam to prevent hydrolysis by class A β -lactamases.

<u>4-Extended spectrum penicillins (Anti-pseudomonal penicillins)</u> like Carbenicillin, Mezlocillin, piperacillin and ticarcillir

Their antimicrobial activity extends to include the *Pseudomonas*, Enterobacter and proteus species as gram negative organisms.

They are destroyed by beta lactamases.

Pharmacokinetics:



- Penicillin G is not used orally (acid labile) and is usually given by Intravenous (IV) or intramuscular (IM) injection.
- Penicillin V is more stable in acidic medium and better absorbed from GIT after oral administration.
- > They are short acting ($\underline{t_{1/2}}$ is 30 minutes) which need frequent administration.
- Penicillin G penetrates readily <u>inflamed meninges</u> to enter the CSF compared with normal meninges.
- ➤ Excretion is mainly by the kidney (10% via glomerular filtration & <u>90% by active tubular secretion</u>).

>To prolong the duration of action and reduce the frequency of penicillin G injection, **probenecid** may be given as it <u>blocks renal</u> tubular secretion of penicillin (but rarely used for this purpose).

Long-acting penicillin

➤The <u>repository preparations of penicillin G (e.g.,</u> penicillin G benzathine) are frequently used in clinical practice.

≻These **I.M**. preparations <u>release</u> penicillin G <u>slowly</u> from the area in which it is injected and produces relatively <u>low but persistent concentrations of antibiotic</u> in the blood.

Penicillin G benzathine preparation is given <u>once per</u> <u>month</u> as a prophylaxis in rheumatic fever.

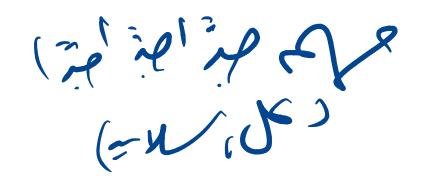
Penicillin procaine is another repository form (long acting) of penicillin but given <u>I.M./12 hours.</u>





Therapeutic uses of penicillin G 5.cocc memori

- 1. Pneumococcal infection: pneumonia and meningitis.
- 2. Streptococcal infection such as pharyngitis caused by β -hemolytic streptococci. This prevents development of acute rheumatic fever, but not glomerulonephritis.
- Penicillin plus aminoglycoside for treatment of streptococcal endocarditis.
- 1. <u>Meningococcal</u> infection: in acute meningitis, but <u>ineffective</u> in meningococcal carrier state or prophylaxis.
- 2. Gonococcal infection, but ceftriaxone is an effective alternative.
- 5- <u>Anaerobic infection</u>: e.g. brain abscess (with metronidazole).
- 6- Syphilis.
- 7-Diphtheria: antitoxin is the only effective treatment, but penicillin
- G eliminates the carrier state.
- 8- <u>Clostridia</u> <u>infections</u>: gas gangrene.
- 9- <u>Anthrax.</u>
- 12- Chemoprophylaxis....



الرقاية الكيمياتي Chemoprophylaxis using Penicillin G and its longacting preparations

Penicillin G is used for Prophylaxis in the following conditions:

- Recurrence of rheumatic fever. Benzathine penicillin G (1.2 million units) given monthly as I.M. injection. In case of hypersensitivity to penicillin, sulfisoxazole or sulfadiazine or macrolides may be alternative.
- 2. Contact persons to patients suffering from syphilis.
- Surgical or dental procedures in cardiac patients with rheumatic valve disease to guard against sub-acute bacterial endocarditis infection (penicillin plus aminoglycoside).

Doses of penicillin G

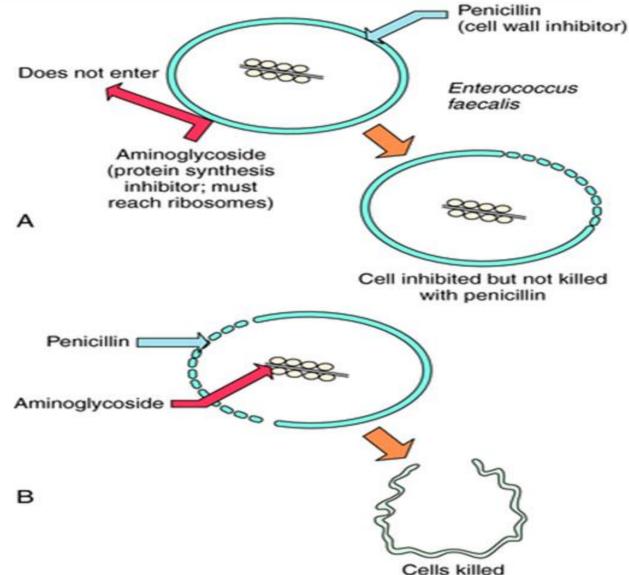
- The dose of penicillin G (4-24 million IU per day) given IV divided into 5 to 6 doses. Each 1 million IU = 0.6 gram of penicillin G.
- Benzathine penicillin G (1.2 million IU) IM (once every 3-4 weeks) is used to prevent recurrence of beta hemolytic streptococcal Infection among patients with rheumatic heart diseases to avoid recurrence of rheumatic fever.
- Benzathine penicillin G (2.4 million IU) IM (once every week for 3-4 weeks) can eradicate syphilis





The combination of penicillin and aminoglycoside

Penicillins and other cell wall inhibitors А facilitate the entry of aminoglycoside into bacterial cells (Synergism) в



II- The penicillinase resistant (anti-staphylococcal) penicillins

Flucloxacillin, Nafcillin, Oxacillin, Cloxacillin, Dicloxacillin, and Methicillin ≻They are resistant to hydrolysis by staphylococcal penicillinases; therefore, their use should be restricted to the treatment of infection caused by staphylococci.

➤ They are less effective against microorganisms susceptible to penicillin G.
 ➤ They have no effect on gram negative bacteria producing penicillinase.

➤Methicillin was withdrawn because of causing interstitial nephritis.
(forever No-one Can destroy me)

Combination of flucloxacillin and amoxicillin are available as oral or injectable preparations.

>Also, combinations of dicloxacillin and ampicillin are available.



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Methicillin resistant microorganisms

Methicillin resistant microorganisms like Methicillin-resistant Staph. aureus (MRSA) is a term applied now to all bacteria which are resistant to all penicillinase resistant penicillins like Methicillin.

MRSA is resistant to most β-lactams because of the presence of mecA, a gene that produces a penicillin binding protein (PBP2a) with low affinity for β-lactam antibiotics

► Vancomycin, linezolid and other drugs is indicated in these conditions although intermediate level of resistance is emerging.





ANTIBIOTICS WITH MRSA COVERAGE

