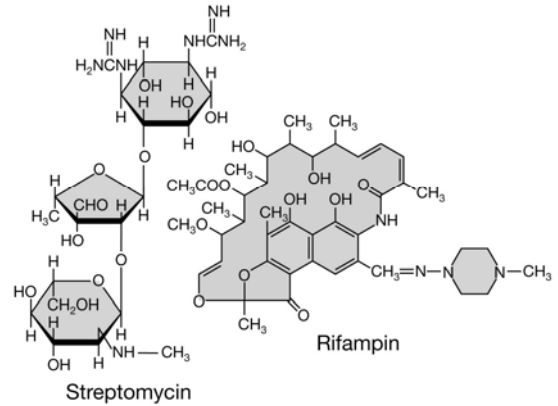


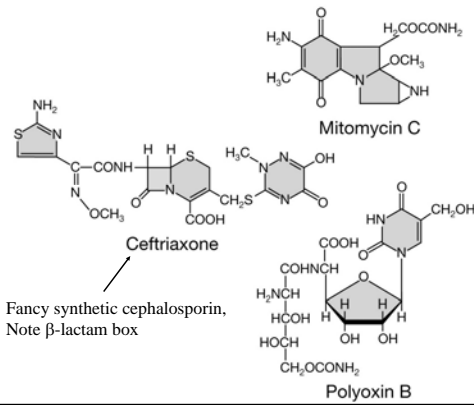
Classification of Antibiotics:

1. Inhibit growth – “stat”
Kill bacterium – “cide”
2. Broad and Narrow spectrum
3. Production Types:
Natural
Synthetic
Semi-synthetic

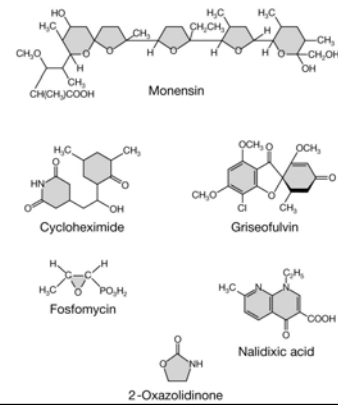
Representative structure



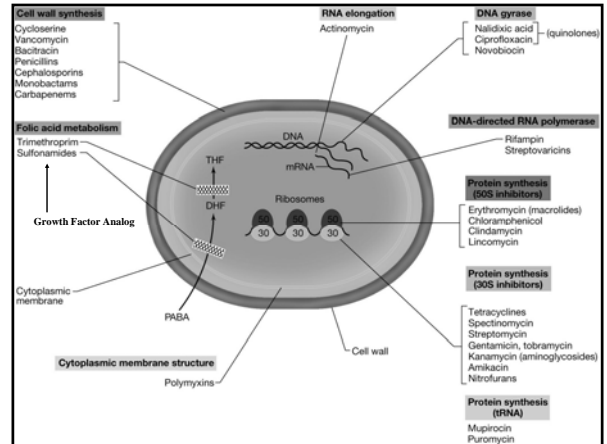
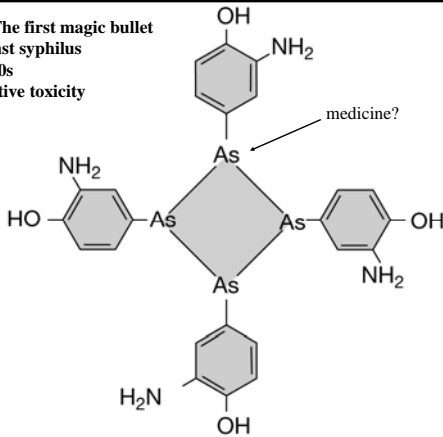
Representative structure



Representative structure



Salvarsan: The first magic bullet
Works against syphilis
Ehrlich, 1900s
Idea of selective toxicity



Antibiotics Affecting Replication, Transcription, & Translation

DNA replication:

Nalidixic Acid & Novobiocin – Inhibits DNA gyrase

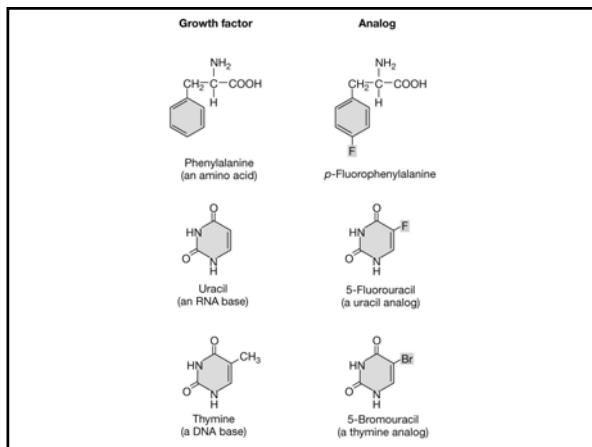
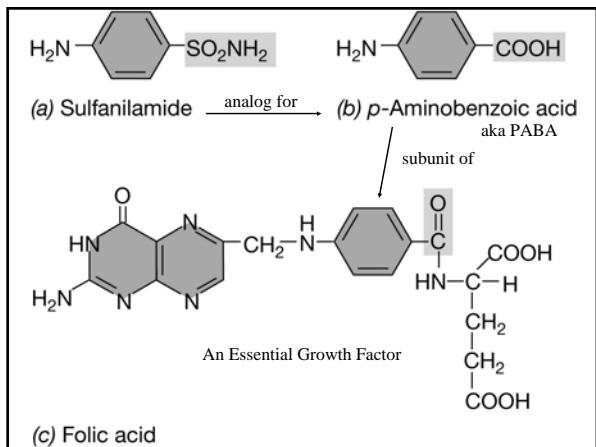
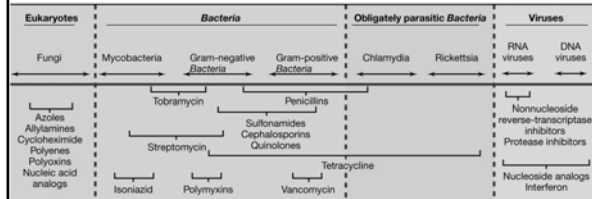
Transcription:

Rifampin – Beta subunit of RNA polymerase
Actinomycin – DNA binding, blocks elongation

Translation:

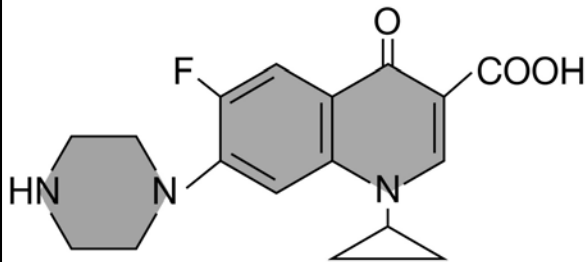
Streptomycin – Blocks initiation on SSU of ribosome
Chloramphenicol – Blocks elongation on LSU via peptide bond
Tetracycline – Blocks elongation SSU
Cycloheximide – Eucarya ribosome specific
Diphtheria Toxin – EF blocker; both Archaea and Eucarya

Antimicrobial spectrum of action for selected chemotherapeutics

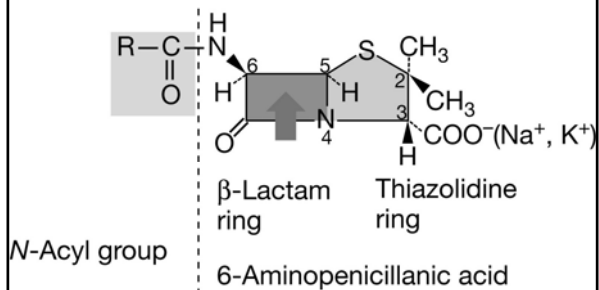


Cipro or ciprofloxacin, a quinolone

Not a growth factor!
Prevents DNA gyrase from supercoiling
Used for anthrax, etc.



How to build a better mouse trap: Penicillin
A β -lactam antibiotic



Inhibits transpeptidation of peptidoglycan chains
Forms the old 1-2-punch with autolysins

Semi's are made to be acid-stable and more broad spectrum

Difference b/t ampicillin and penicillin is only one amino group.

Designation	N-Acyl group
NATURAL PENICILLIN Benzylpenicillin (penicillin G) Gram-positive activity β-lactamase-sensitive	
SEMISYNTHETIC PENICILLINS	
Methicillin acid-stable, β-lactamase-resistant	
Oxacillin acid-stable, β-lactamase-resistant	
Ampicillin broadened spectrum of activity (especially against gram-negative bacteria), acid-stable, β-lactamase-resistant	
Carbenicillin broadened spectrum of activity (especially against Pseudomonas aeruginosa), acid-stable but ineffective orally, β-lactamase-sensitive	

Figure 20.19 The structures of some important penicillins.

N-Acetyltransferase — R-plasmid born resistance

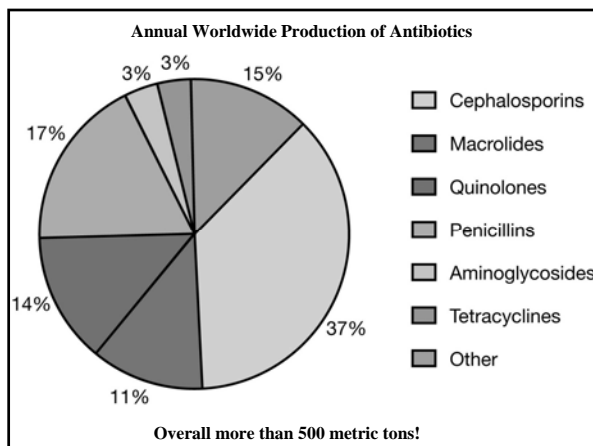
Kanamycin, an aminoglycoside
Inhibits protein synthesis on small subunit rRNA
Considered a reserve antibiotic

Macrolide ring

Erythromycin, a macrolide
Inhibits protein synthesis on large subunit rRNA
Often works when allergic to β-lactams

Tetracyclines
Inhibits protein synthesis on small subunit rRNA
Along with β-lactams make up the majority used

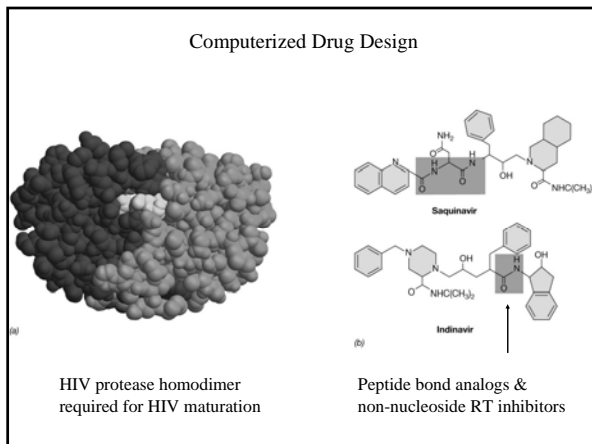
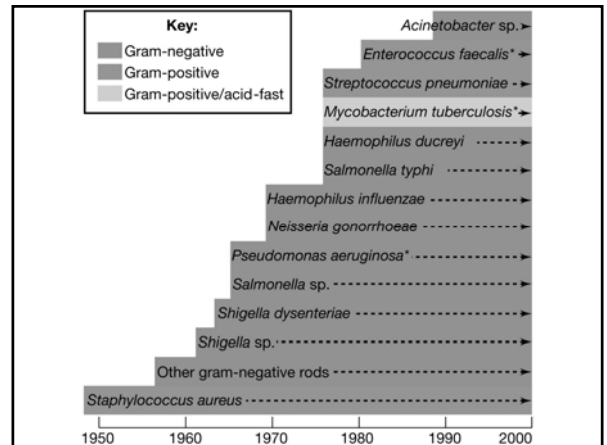
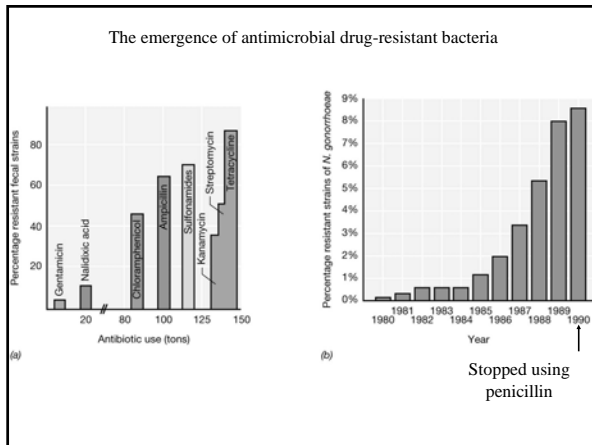
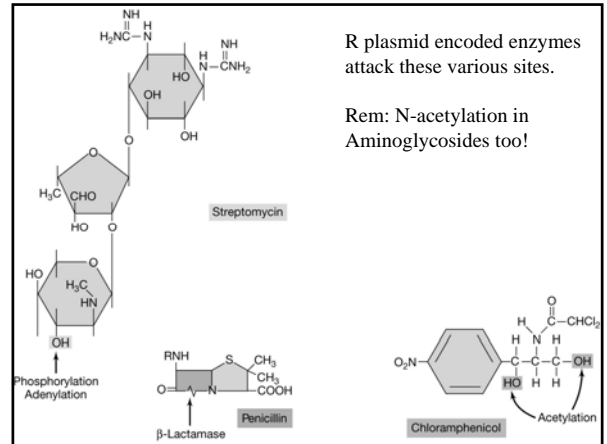
Tetracycline analog	R ₁	R ₂	R ₃	R ₄
Tetracycline	H	OH	CH ₃	H
7-Chlortetracycline (aureomycin)	H	OH	CH ₃	Cl
5-Oxytetracycline (terracycline)	OH	OH	CH ₃	H



- Mechanisms of Antibiotic Resistance**
- Lacks structure antibiotic inhibits:
Mycoplasmas lack a typical cell wall
 - Impermeable to the antibiotic:
Gram - bacteria impermeable to penicillin G
 - Alteration of antibiotic:
β-lactamase degrades antibiotic e.g., springs open the mouse trap
 - Modifies the target of the antibiotic
 - Genetically modifies the pathway that the antibiotic affects
 - Efflux of the antibiotic:
Tetracycline gets pumped back out of the cell

TABLE 20.7 Mechanisms of bacterial resistance to antibiotics

Resistance mechanism	Antibiotic example	Genetic basis of resistance	Mechanism present in:
Reduced permeability	Penicillins	Chromosomal	<i>Pseudomonas aeruginosa</i> Enteric Bacteria
Inactivation of antibiotic (for example, penicillinase; modifying enzymes methylases, acetylases, and phosphorylases; and others)	Penicillins	Plasmid and chromosomal	<i>Staphylococcus aureus</i> Enteric Bacteria
Alteration of target (for example, RNA polymerase, rifamycin, ribosome, erythromycin, and streptomycin; DNA gyrase, quinolones)	Chloramphenicol	Plasmid and chromosomal	<i>Neisseria gonorrhoeae</i> <i>Staphylococcus aureus</i> Enteric Bacteria
Development of resistant biochemical pathway	Aminoglycosides	Plasmid	<i>Staphylococcus aureus</i> Enteric Bacteria
Efflux (pumping out of cell)	Erythromycin Rifamycin Streptomycin Norfloxacin	Chromosomal	<i>Staphylococcus aureus</i> Enteric Bacteria
	Sulfonamides	Chromosomal	<i>Staphylococcus aureus</i> Enteric Bacteria
	Tetracyclines	Plasmid	Enteric Bacteria
	Chloramphenicol	Chromosomal	<i>Staphylococcus aureus</i> <i>Bacillus subtilis</i>



Microbial Sources of Antibiotics

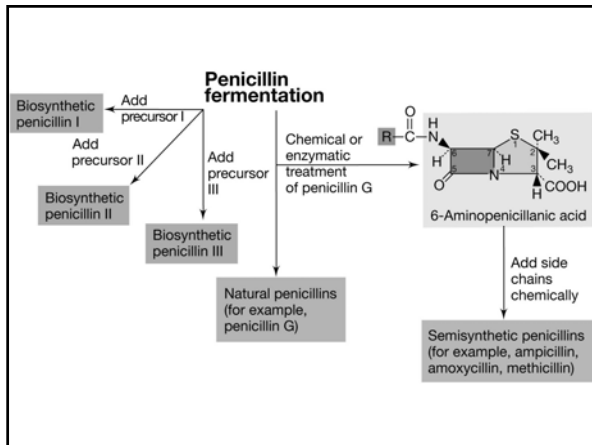
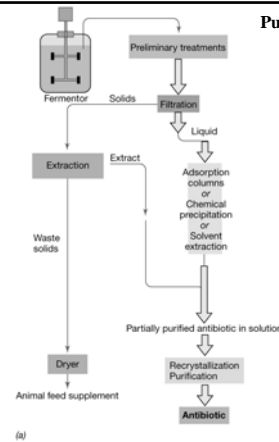
Microorganism	Antibiotic
Bacteria:	
<i>Streptomyces</i> spp.	chloramphenicol erythromycin kanamycin rifampin streptomycin tetracyclines
<i>Bacillus</i> spp.	bacitracin polymyxin
Fungi:	
<i>Penicillium</i> spp.	penicillin
<i>Cephalosporium</i> spp.	cephalosporins

Production of Antibiotics:

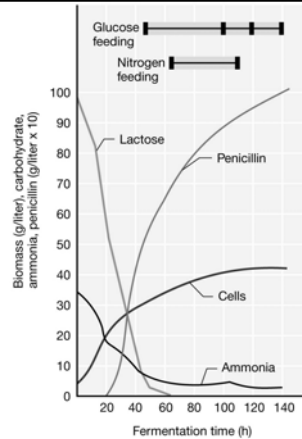
Secondary Metabolites produced near the end of a bacterium or fungus life cycle:

1. Formed @ end of stationary phase of growth
2. Not essential for growth or viability
3. Formation depends upon the media, possible over production

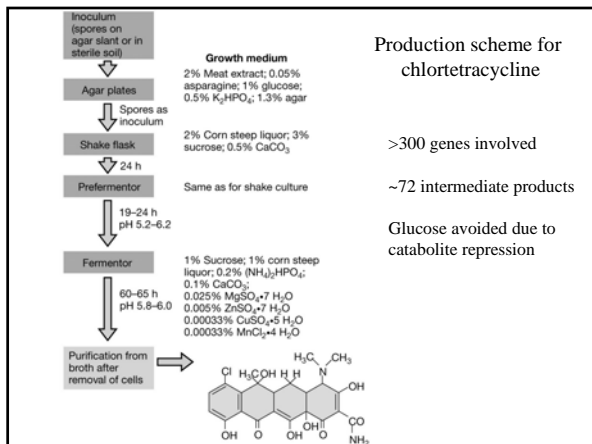
Purification of an antibiotic



Kinetics of penicillin fermentation



Production scheme for chlortetracycline



>300 genes involved
~72 intermediate products
Glucose avoided due to catabolite repression