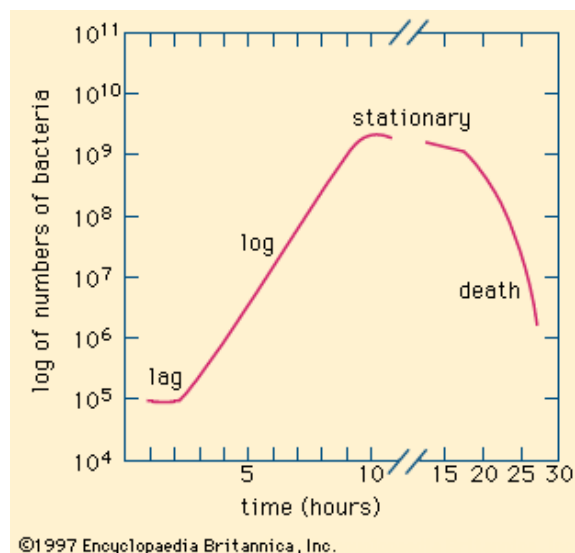


Microbial Growth Control

Antibiotics
Bacteriocins
Bdellovibrio – a bacteria eater
Bacterial cannibalism

Effect of Antimicrobial Agents on Growth

Growth Curve of Bacteria



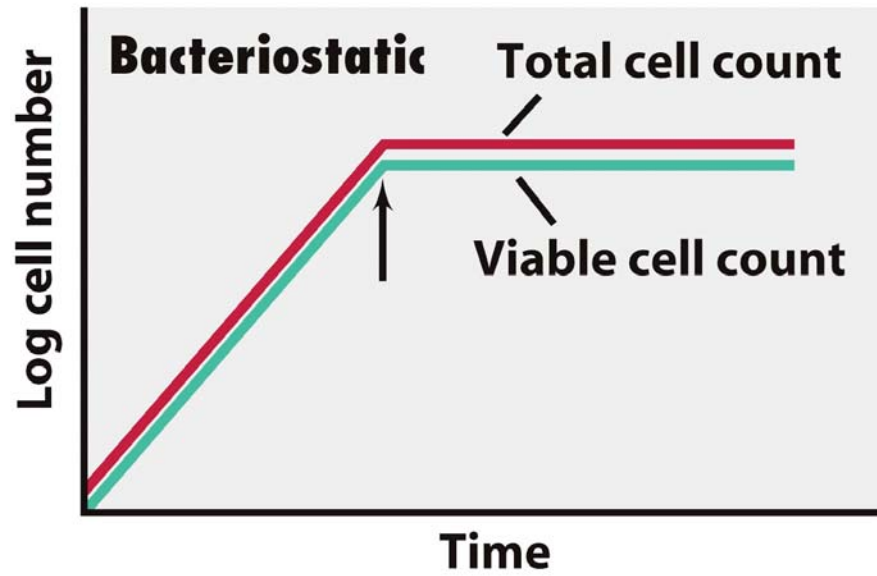


Figure 20-9a Brock Biology of Microorganisms 11/e
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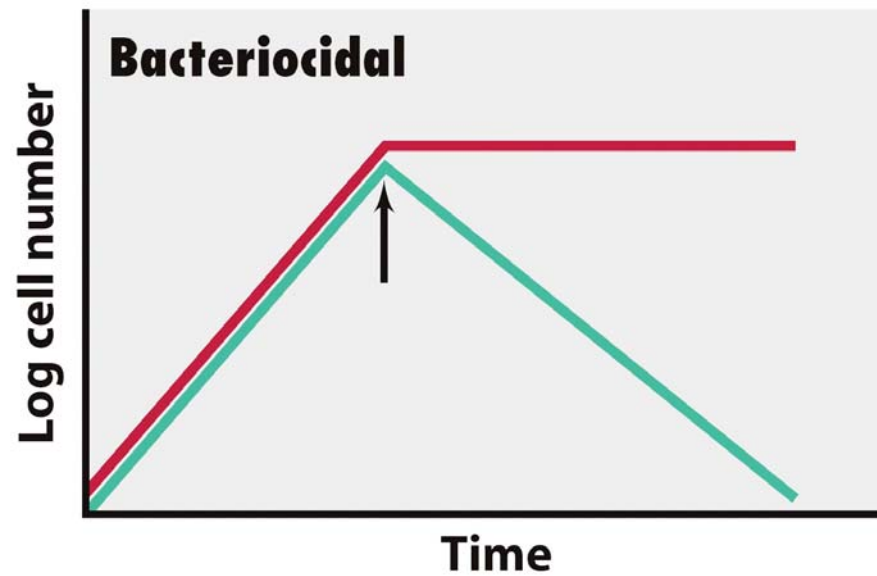


Figure 20-9b Brock Biology of Microorganisms 11/e
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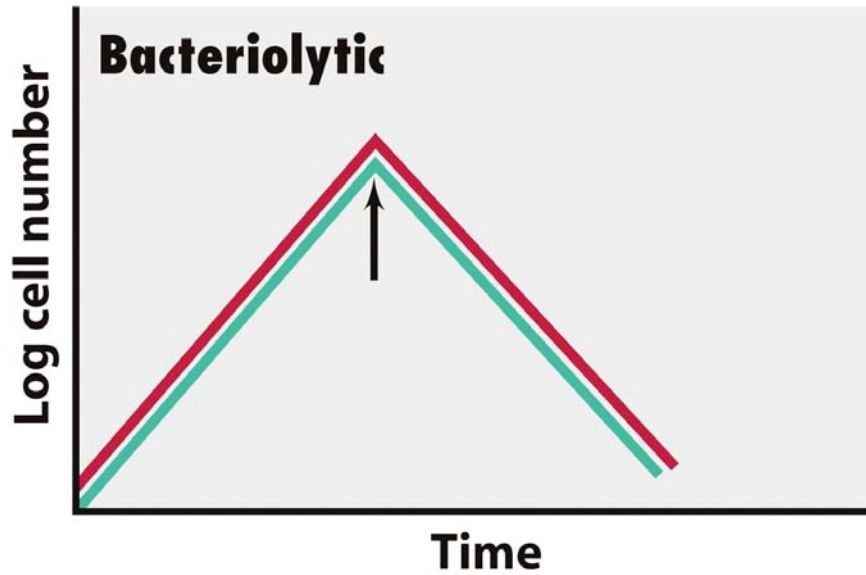


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Measuring Antimicrobial Activity

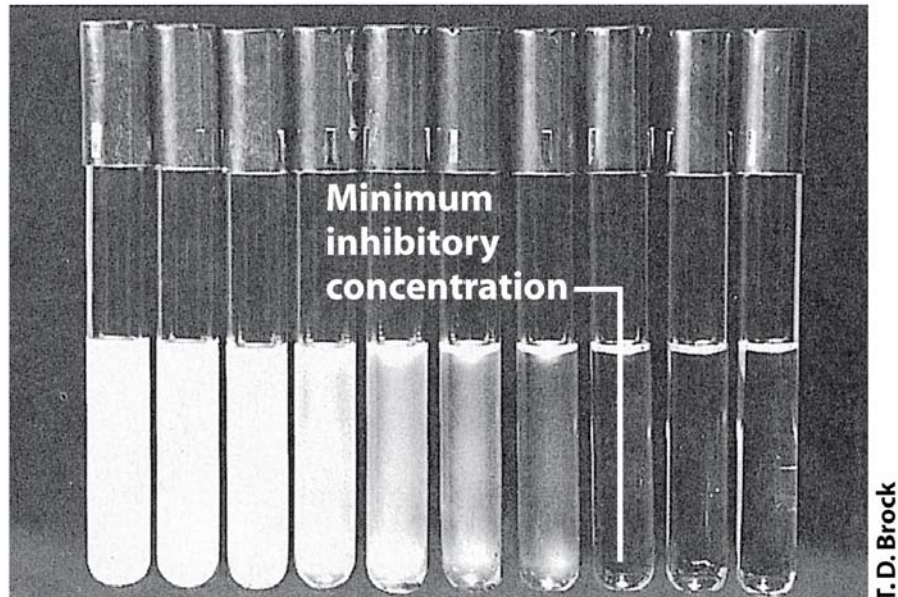


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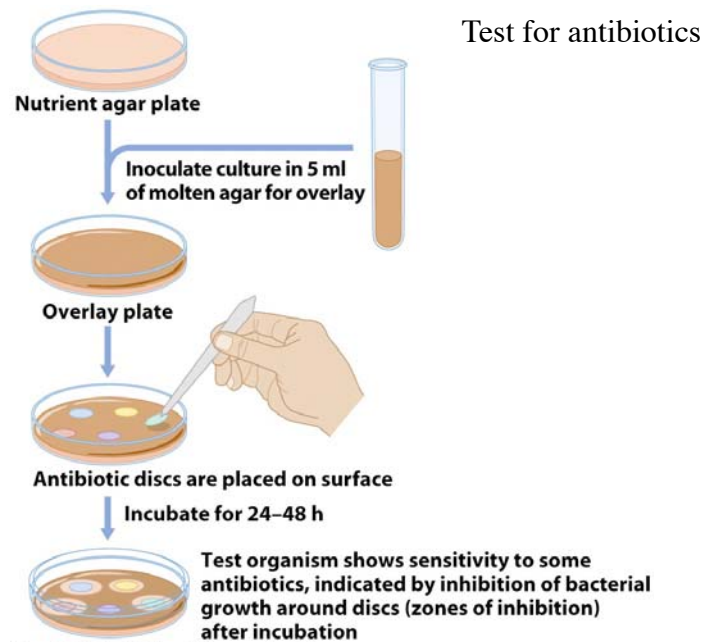
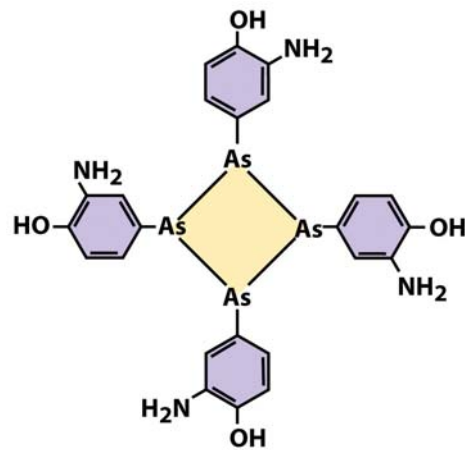


Figure 20-11 Brock Biology of Microorganisms 11/e
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Salvarsan:
Treatment of Syphilis
In the early 1910
“compound 606”

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Targets of antibiotics

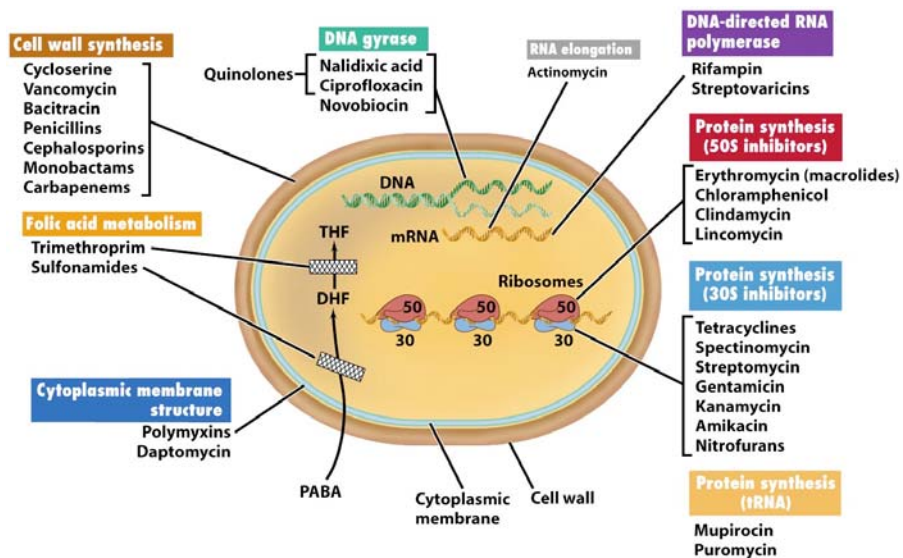


Figure 20-14 Brock Biology of Microorganisms 11/e
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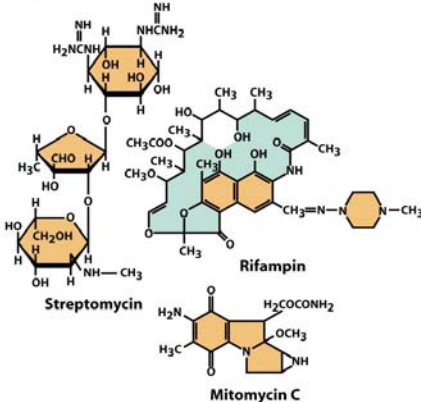
Antibiotic classification	Subclassification	Example	Representative structure
I. Carbohydrate-containing compounds	Pure sugars	Nojirimycin	
	Aminoglycosides	Streptomycin	
	Orthosomycins	Everninomycin	
	N-Glycosides	Streptothricin	
	C-Glycosides	Vancomycin	
II. Macrocyclic lactones	Glycolipids	Moenomycin	
	Macrolide antibiotics	Erythromycin	
	Polyene antibiotics	Candididin	
	Ansamycins	Rifampin	
III. Quinones and related compounds	Macrotetrolides	Tetranactin	
	Tetracyclines	Tetracycline	
	Anthracyclines	Adriamycin	
	Naphthoquinones	Mitomycin C	
	Benzoquinones	Actinorhodin	
		Mitomycin C	

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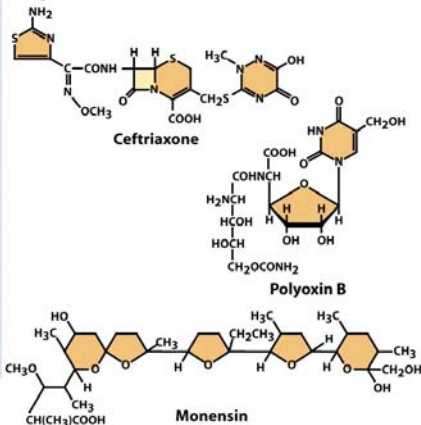
Antibiotic classification	Subclassification	Example	Representative structure
IV. Amino acid and peptide analogs	Amino acid derivatives	Cycloserine	
	β-Lactam antibiotics	Penicillin, ceftriaxone	
	Peptide antibiotics	Bacitracin	
	Chromopeptides	Actinomycin	
	Depsideptides	Valinomycin	
V. Heterocyclic compounds containing nitrogen	Chelate-forming peptides	Bleomycin	
V. Heterocyclic compounds containing nitrogen	Nucleoside antibiotics	Polyoxins	
VI. Heterocyclic compounds containing oxygen			
	Polyether antibiotics	Monensin	

Figure 20-13 part 2 Brock Biology of Microorganisms 11/e
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Antibiotic classification	Subclassification	Example
VII. Alicyclic derivatives	Cycloalkane derivatives Steroid antibiotics	Cycloheximide Fusidic acid
VIII. Aromatic compounds	Benzene derivatives Condensed aromatics Aromatic ether	Chloramphenicol Griseofulvin Novobiocin
IX. Aliphatic compounds	Compounds containing phosphorus	Fosfomycin

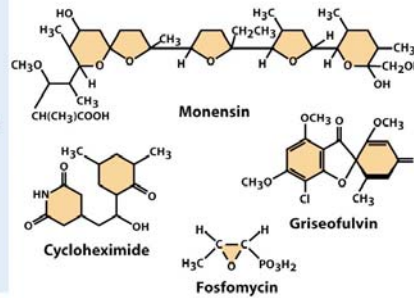
Representative structure

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Antibiotic classification	Subclassification	Example
X. Quinolone compounds	4-Quinolone Fluoro-4-quinolones	Nalidixic acid Ciprofloxacin
XI. Oxazolidinone	Cyclic lactone	2-Oxazolidinone

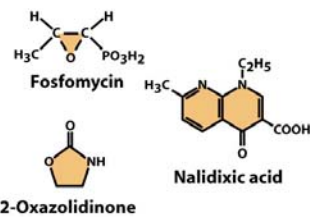
Representative structure

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Wirkspektren

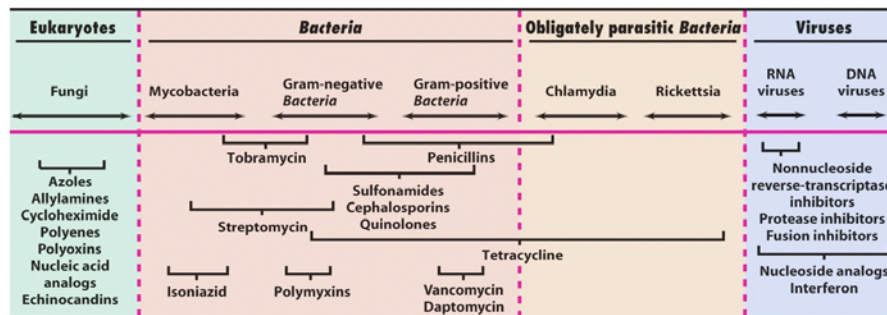


Figure 20-15 Brock Biology of Microorganisms 11/e
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Annual worldwide production and use of antibiotics

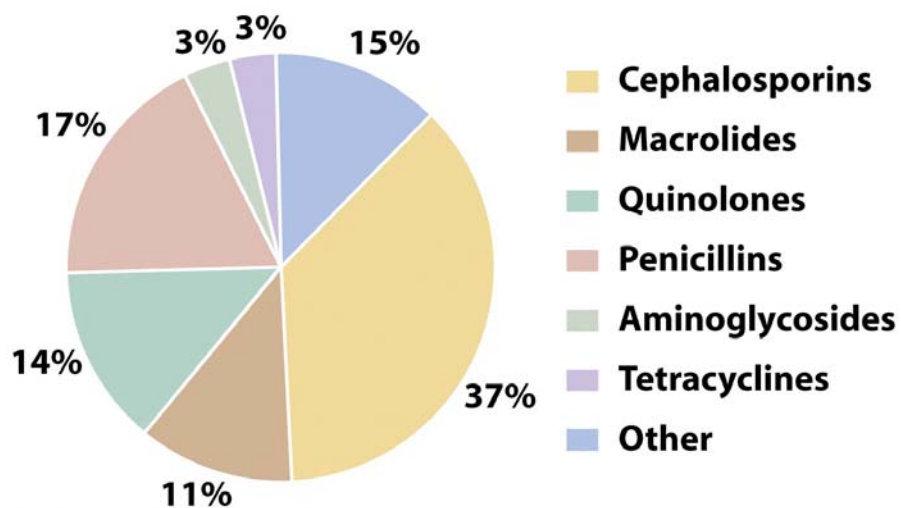
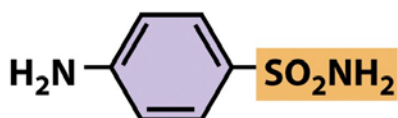


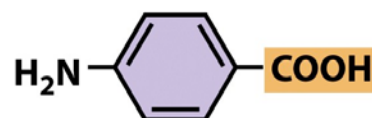
Figure 20-16 Brock Biology of Microorganisms 11/e
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Sulfa Drugs

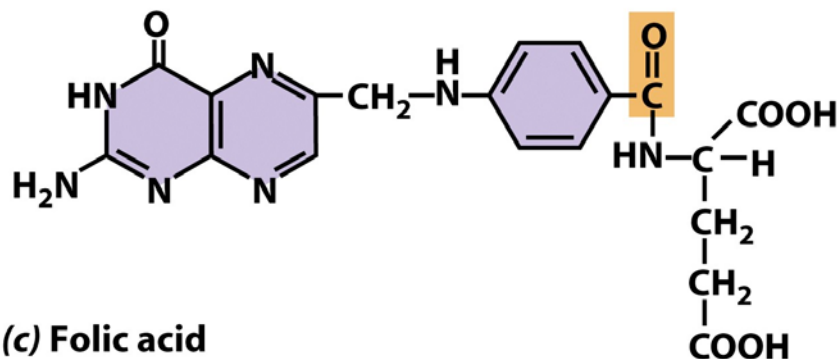
Sulfanilamid is an analog of paba, a precursor of the growth factor folic acid



(a) Sulfanilamide



(b) *p*-Aminobenzoic acid



(c) Folic acid

Figure 20-17 Brock Biology of Microorganisms 11/e
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Nucleic Acid Analogs

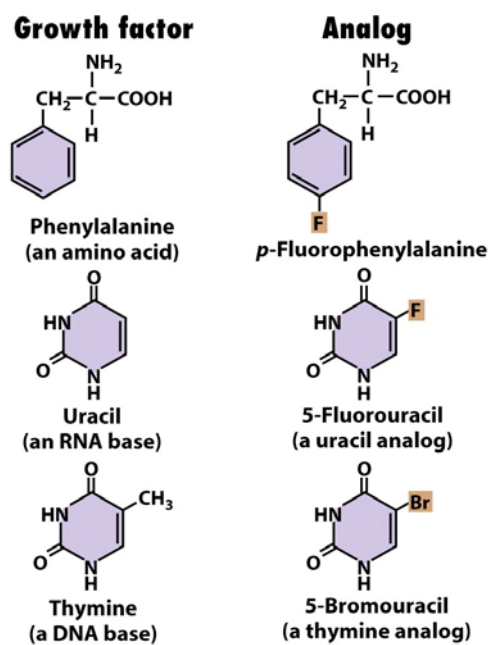


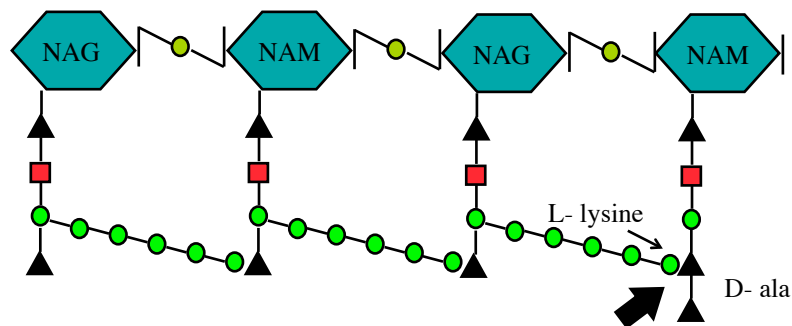
Figure 20-18 Brock Biology of Microorganisms 11/e
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Naturally Occurring Antimicrobial Drugs: Antibiotics

β -Lactam Antibiotics: Penicillins and Cephalosporins

Inhibitors of bacterial cell wall synthesis

Cell wall Attachment of new wall unit to growing peptidoglycan



Beta-lactams

Bind to and inhibit enzymes which catalyse this link

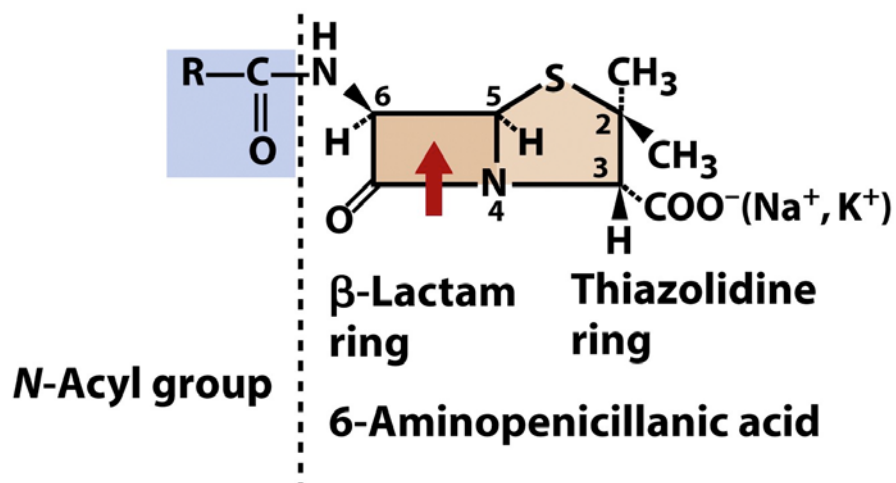


Figure 20-20 part 1 Brock Biology of Microorganisms 11/e
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Alexander Fleming in 1928

The β -lactam compounds, including the penicillins and the cephalosporins, are the most important clinical antibiotics. These antibiotics target cell wall synthesis in *Bacteria*. They have low host toxicity and a broad spectrum of activity.

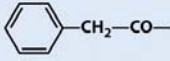
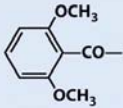
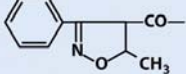
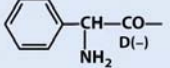
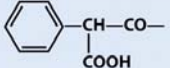
Designation	N-Acyl group
NATURAL PENICILLIN Benzympenicillin (penicillin G) Gram-positive activity β -lactamase-sensitive	
SEMISYNTHETIC PENICILLINS	
Methicillin acid-stable, β -lactamase-resistant	
Oxacillin acid-stable, β -lactamase-resistant	
Ampicillin broadened spectrum of activity (especially against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

Figure 20-20 part 2 Brock Biology of Microorganisms 11/e
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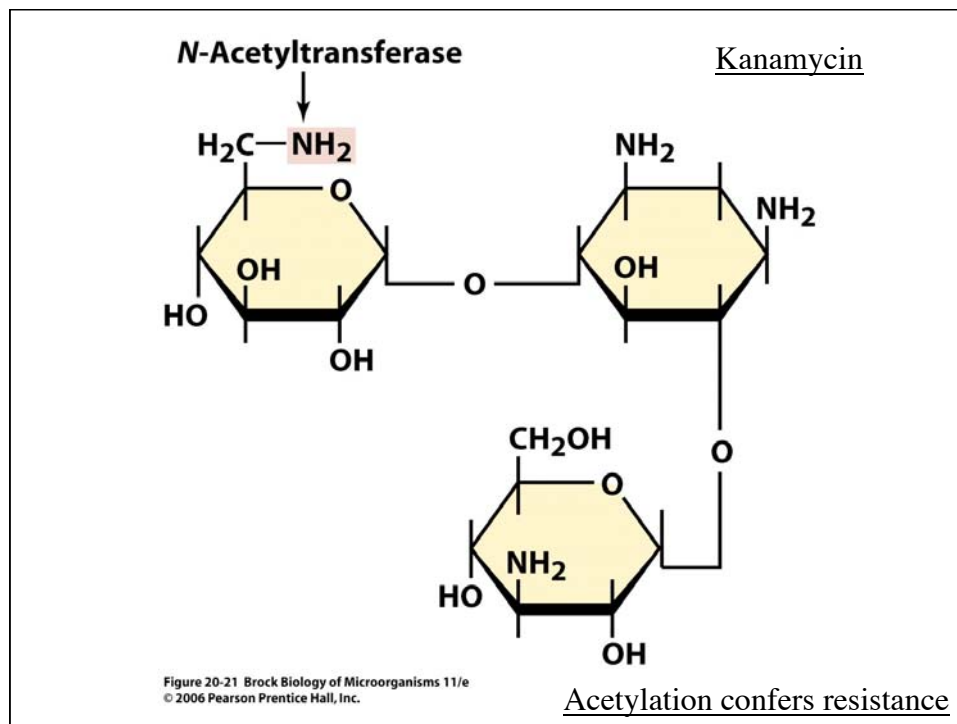
Antibiotics from Prokaryotes

Aminoglycoside Antibiotics

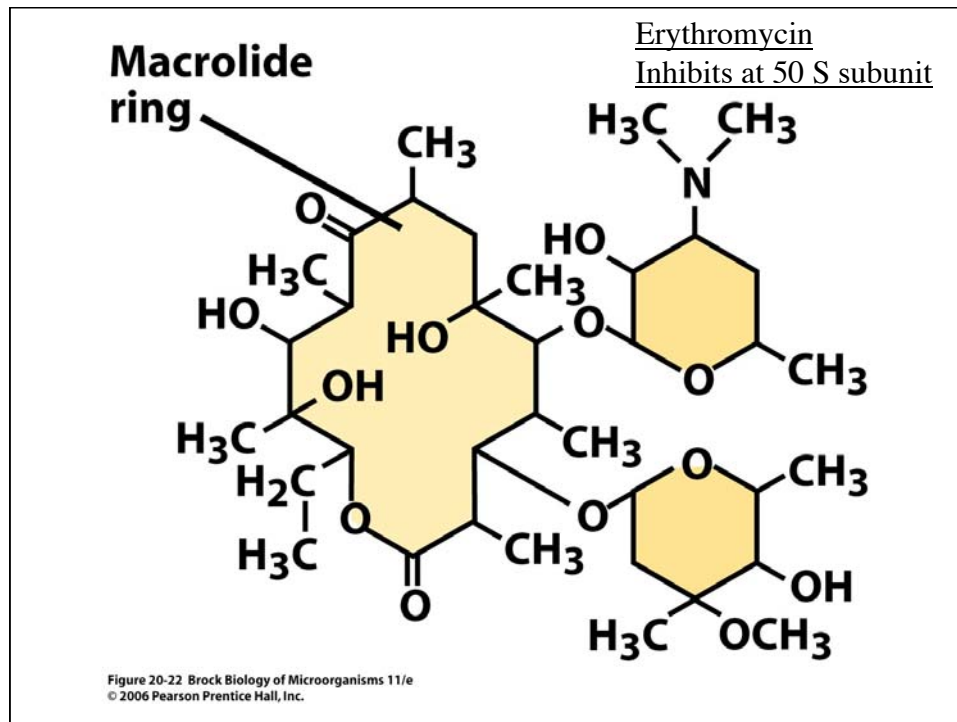
Amino sugars bonded by glycosidic linkages

Inhibit 30S subunit

Against gram-negative bacteria

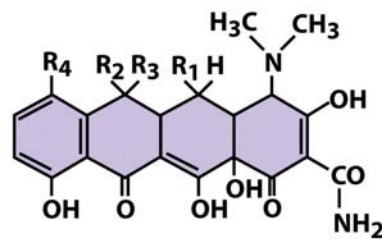


Macrolide Antibiotics



Tetracyclines

Tetracyclins interfere with
30S ribosomal subunit



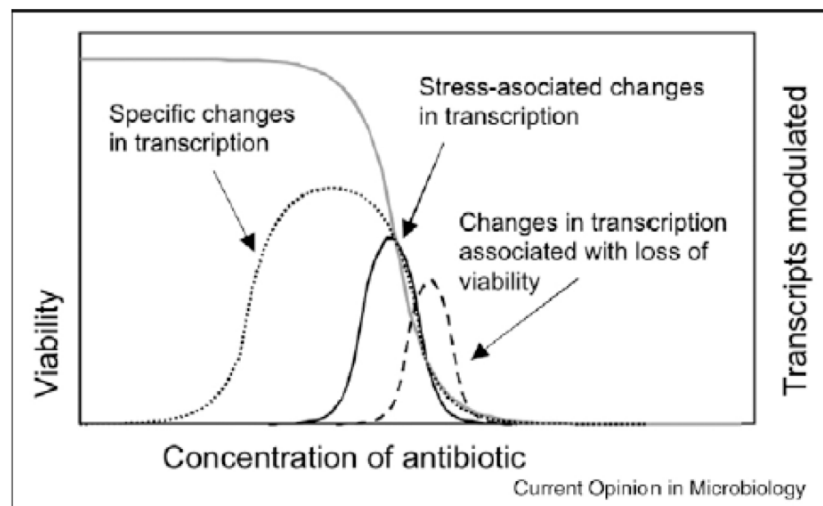
Tetracycline analog	R₁	R₂	R₃	R₄
Tetracycline	H	OH	CH ₃	H
7-Chlortetracycline (aureomycin)	H	OH	CH ₃	Cl
5-Oxytetracycline (terramycin)	OH	OH	CH ₃	H

Figure 20-23 Brock Biology of Microorganisms 11/e
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The aminoglycosides, macrolides, and tetracycline antibiotics are structurally complex molecules produced by *Bacteria* and are active against other *Bacteria*. All of these work by interfering with protein synthesis.

A different View on Antibiotics

Antibiotics in Nature: Signalling molecules at low concentration



Resistance Mechanisms

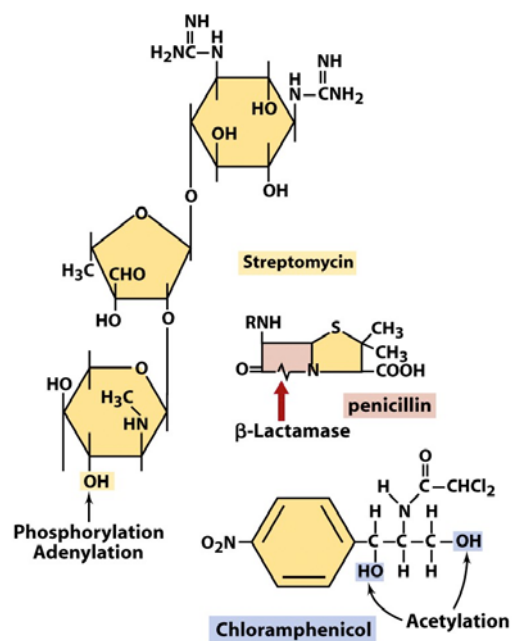
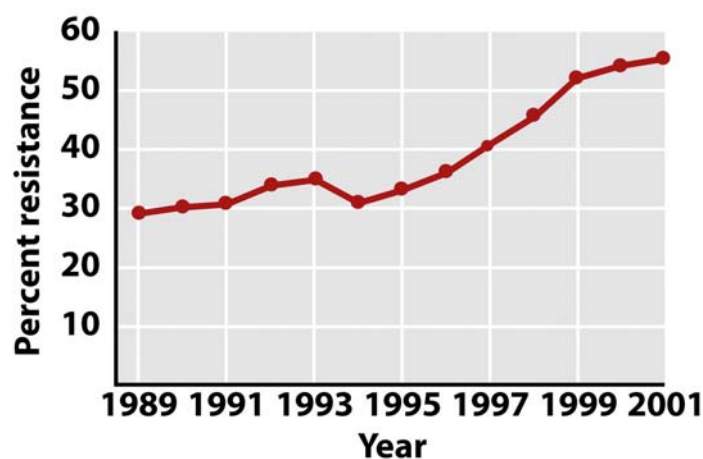


Figure 20-25 Brock Biology of Microorganisms 11/e
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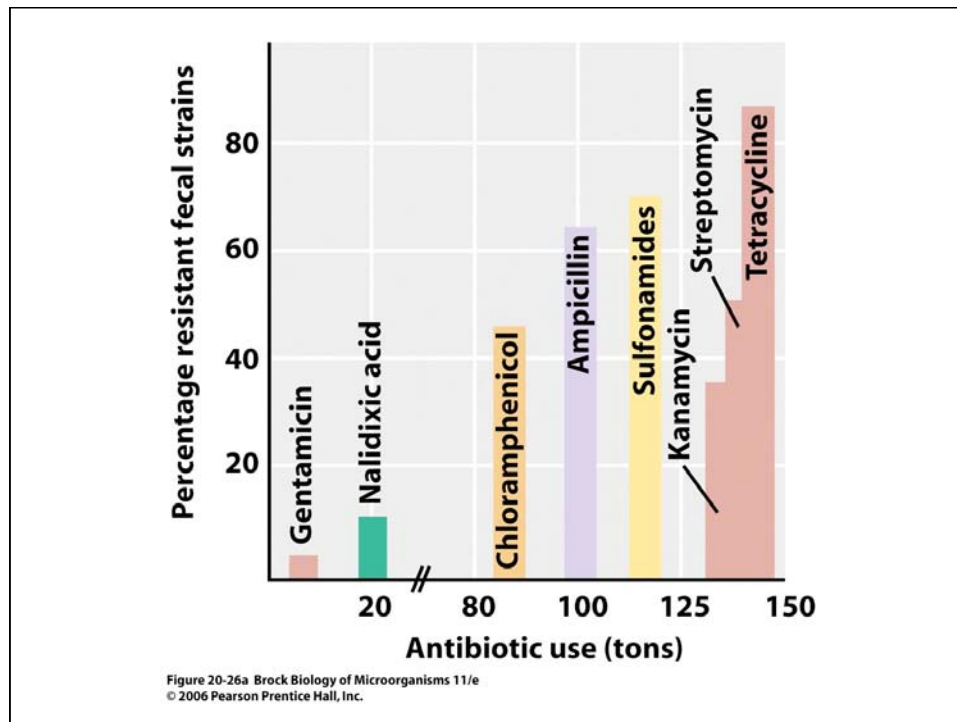
Table 20.7 Mechanisms of bacterial resistance to antibiotics			
Resistance mechanism	Antibiotic example	Genetic basis of resistance	Mechanism present in:
Reduced permeability	Penicillins	Chromosomal	<i>Pseudomonas aeruginosa</i> Enteric Bacteria
Inactivation of antibiotic (for example, penicillinase; modifying enzymes methylases, acetylases, and phosphorylases; and others)	Penicillins	Plasmid and chromosomal	<i>Staphylococcus aureus</i> Enteric Bacteria
	Chloramphenicol	Plasmid and chromosomal	<i>Neisseria gonorrhoeae</i> <i>Staphylococcus aureus</i> Enteric Bacteria
Alteration of target (for example, RNA polymerase, rifamycin; ribosome, erythromycin, and streptomycin; DNA gyrase, quinolones)	Aminoglycosides	Plasmid	<i>Staphylococcus aureus</i>
	Erythromycin	Chromosomal	<i>Staphylococcus aureus</i>
	Rifamycin		Enteric Bacteria
	Streptomycin		Enteric Bacteria
Development of resistant biochemical pathway	Norfloxacin		Enteric Bacteria
	Sulfonamides	Chromosomal	<i>Staphylococcus aureus</i> Enteric Bacteria
Efflux (pumping out of cell)	Tetracyclines	Plasmid	<i>Staphylococcus aureus</i>
	Chloramphenicol	Chromosomal	Enteric Bacteria
	Erythromycin	Chromosomal	<i>Staphylococcus aureus</i> <i>Bacillus subtilis</i> <i>Staphylococcus</i> spp.

Table 20-7 Brock Biology of Microorganisms 11/e
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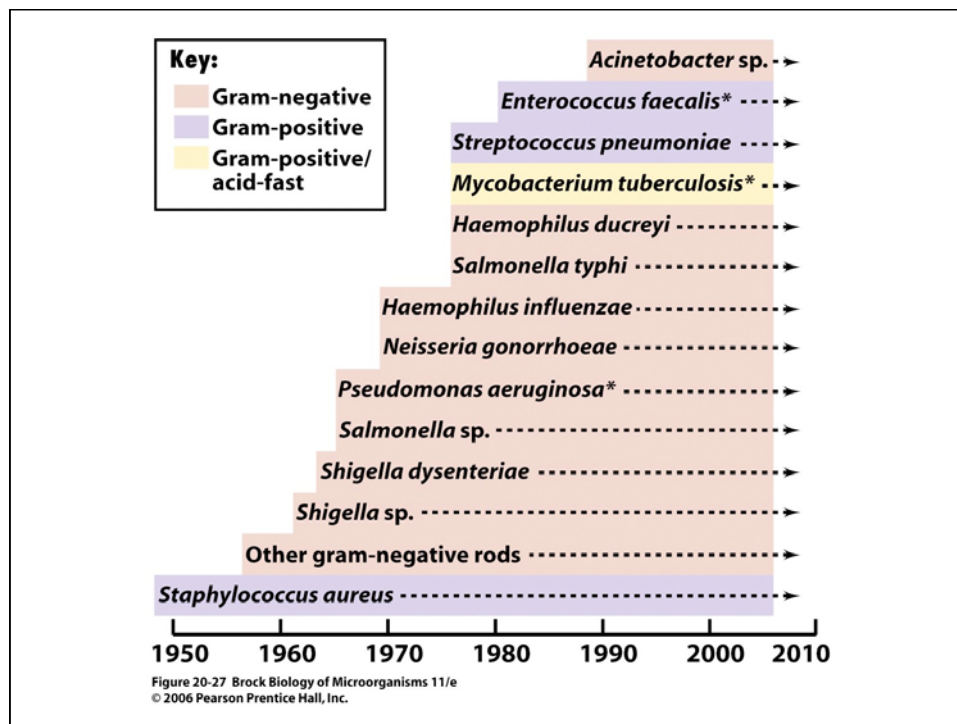
Methicillin-resistant *Staphylococcus aureus*
nosocomial infections among intensive care unit patients
from 1989-2001



Sidebar 20-1 Brock Biology of Microorganisms 11/e
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Antibiotic-Resistant Pathogens

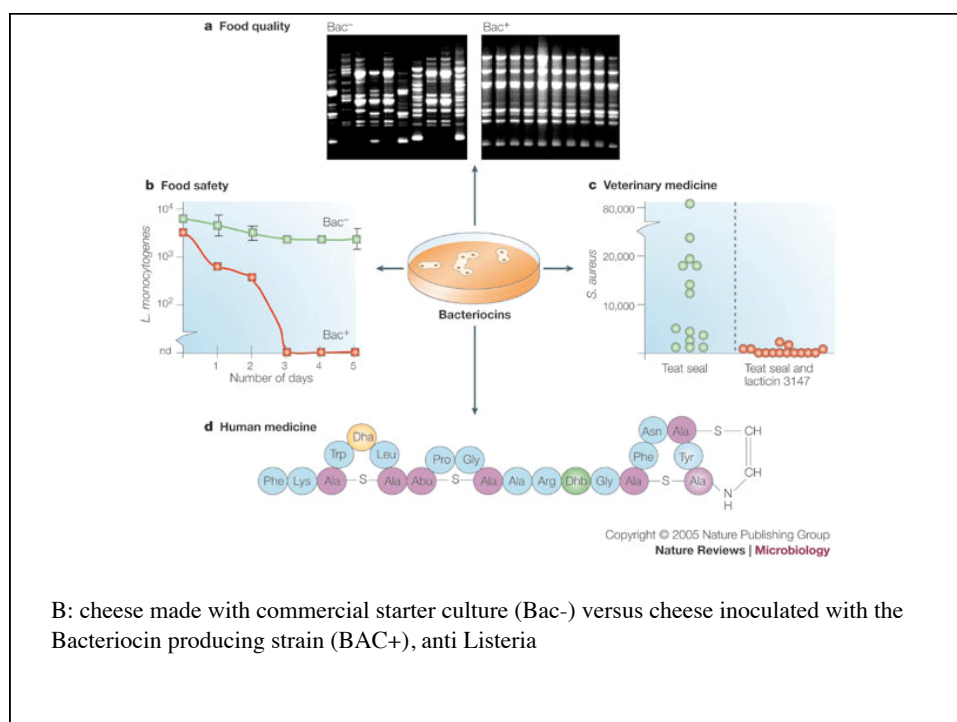
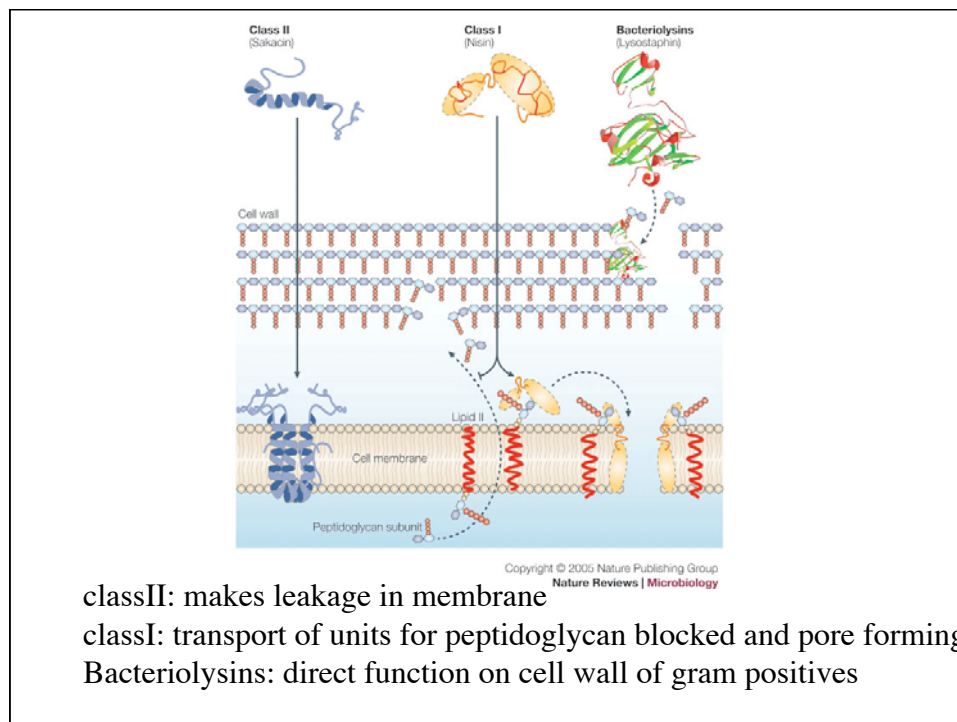


Bacteriocins

bacterially produced, small heat-stable peptides,
active against other bacteria
producer has specific immunity mechanism
narrow or broad target spectrum

many are produced by food-grade lactic acid bacteria
helps to direct or prevent the development of specific

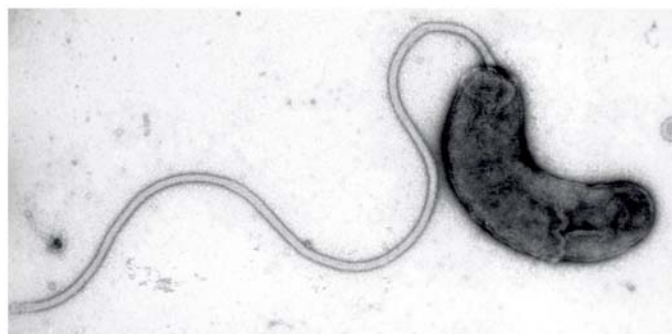
bacterial species in food.



Bdellovibrio: a bacterial killer

Bdellovibrio:

obligate aerobe,
highly motile,
energy from the oxidation of AA and Acetate and compounds of other bacteria
replicates in periplasmic space
spherical structure called bdelloplast
attacks only gram- bacteria
widespread in soil and water, including marine environments
can be isolated like viruses (plaque assay with growing plaque)



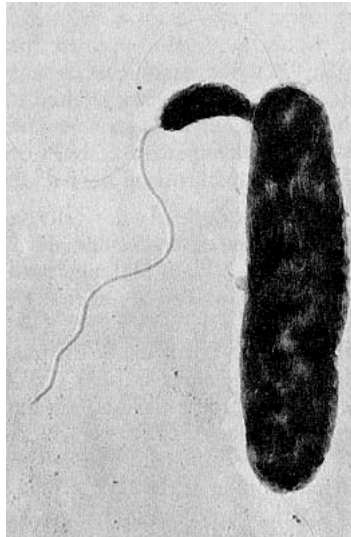
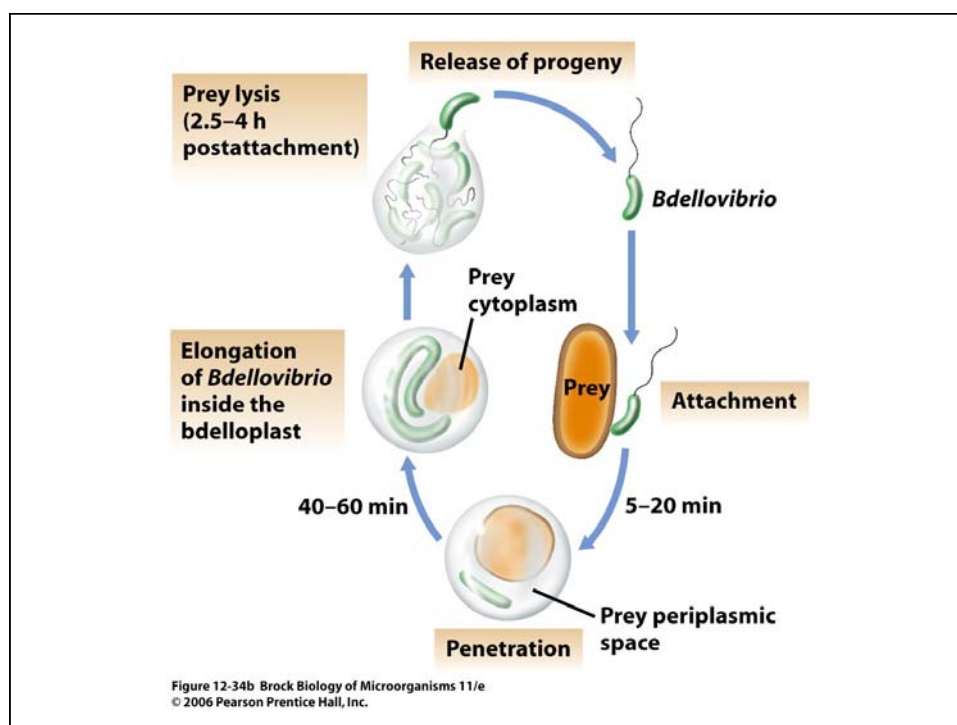
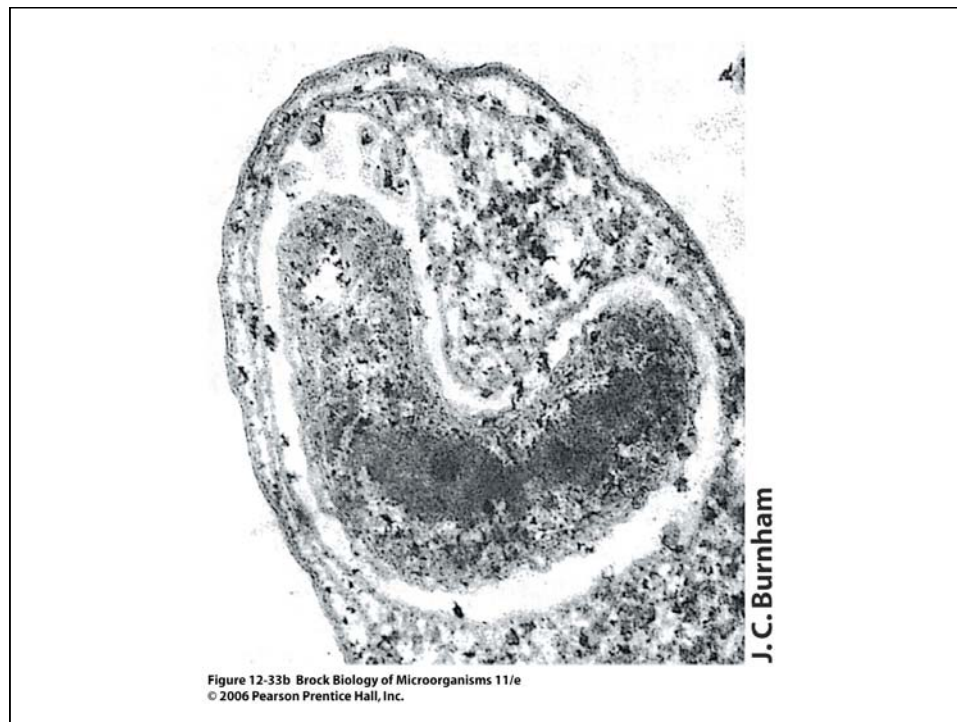


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J. C. Burnham





Bacterial cannibalism and fratricide

