Comparing Bacteria, Archaea and Eucarya

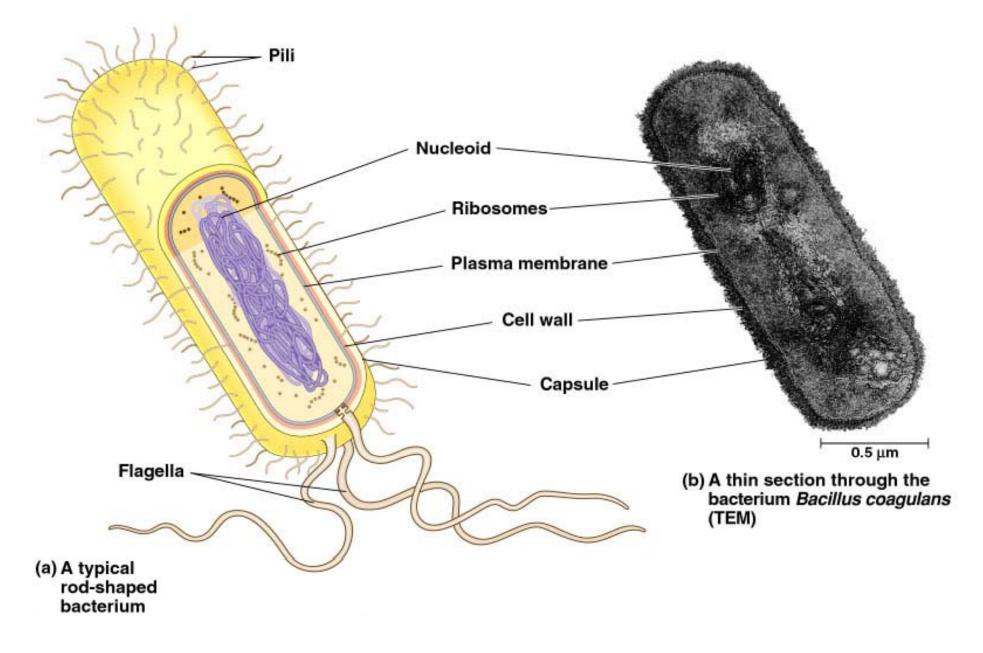
Basic unit of living organisms is the cell; the smallest unit capable of life.

"Features" found in all cells:

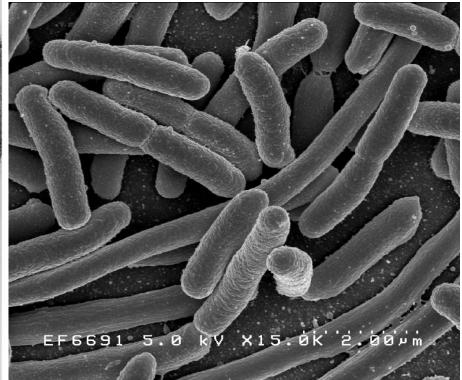
- Ribosomes
- Cell Membrane
- Genetic Material
- Cytoplasm

- ATP Energy
- External Stimuli
- Regulate Flow
- Reproduce

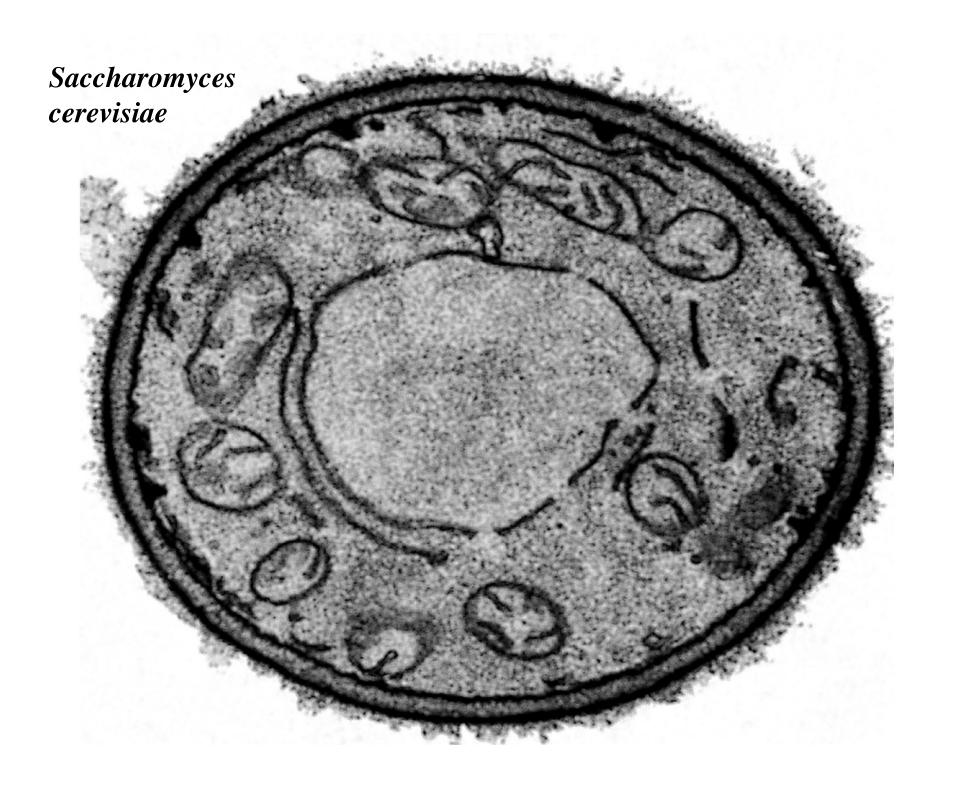
A Bacterial Cell



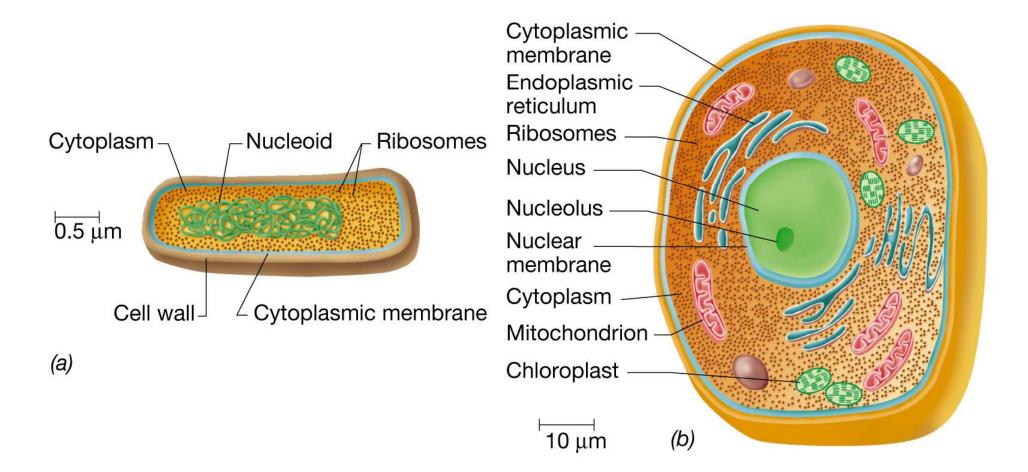




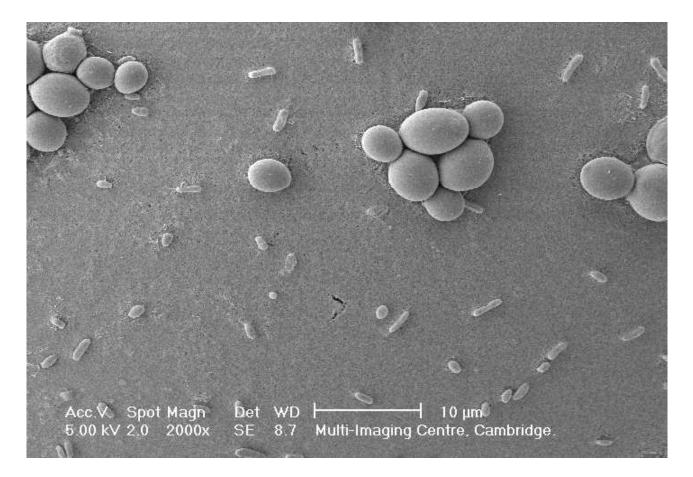
Escherichia coli

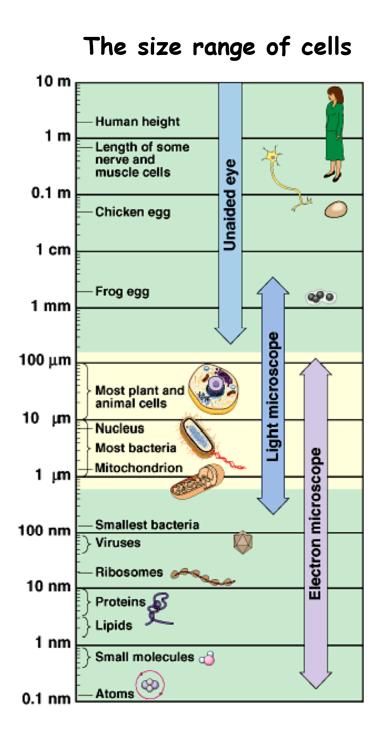


Elements of cellular structure



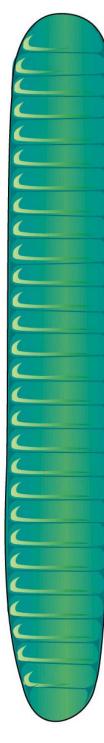
E. coli and S. cerevisiae





Oscillatoria (a cyanobacterium) $8 \times 50 \ \mu m$

Size relationship among Bacteria



Bacillus megaterium $1.5 \times 4 \ \mu m$



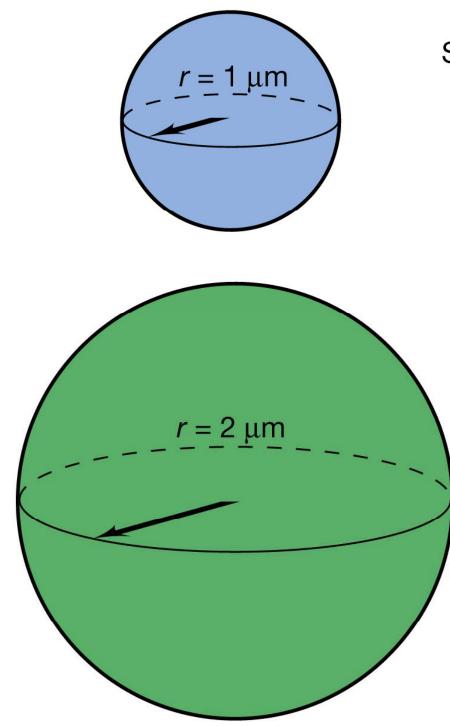
Escherichia coli $1 \times 3 \ \mu m$



Streptococcus pneumoniae 0.8 μm diameter



Haemophilus influenzae $0.25 \times 1.2 \ \mu m$



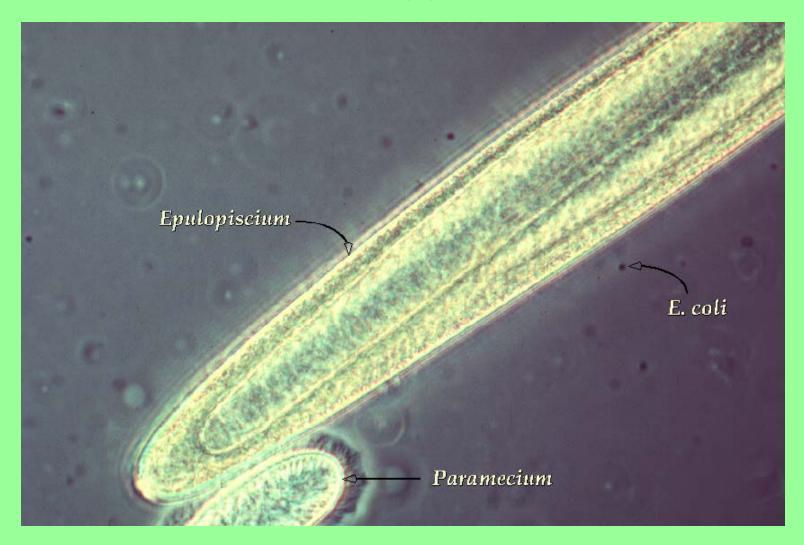
Surface area $(4\pi r^2) = 12.6 \ \mu m^2$ Volume $(\frac{4}{3}\pi r^3) = 4.2 \ \mu m^3$

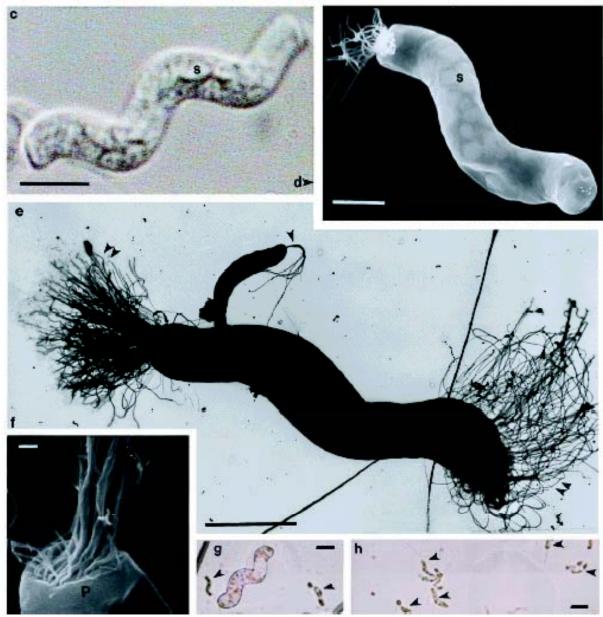
 $\frac{\text{Surface}}{\text{Volume}} = 3$

Surface area = 50.3 μ m² Volume = 33.5 μ m³

 $\frac{\text{Surface}}{\text{Volume}} = 1.5$

A Million times bigger than E. coli!





Titanospirillum velox

Up to 40 µm long

Fto: 1. (a) Mat surface at the Ebro Delta field site (3) showing lack of standing water. (Bar -10 cm.) (b) Two spirilla cells (S, sulfor globale) shown by differential interference contrast (Nomarski). (Bar $-5 \mu \text{m.}$) (c) Phase contrast microscopy of live spirillum cells. (Bar $-5 \mu \text{m.}$) (d) Bipolar lophotrichous large spirillum in which only one pole has retained flagella. Sulfur globales are visible through the cell wall (scanning electron micrograph). (Bar $-5 \mu \text{m.}$) (e) Negative-stain transmission electron micrograph of an entire bipolar lophotrichous large spirillum showing flagella "braids" (double arrowheads) compared with standard-sized spirilla (single arrowhead). (Bar $-5 \mu \text{m.}$) (f) This scanning-electron micrograph of a cell terminous shows one vaulted end with residual flagella. The indentation coated by the polar organelle (P; see Fig. 2) is implied. (Bar $-0.5 \mu \text{m.}$) (g) This Gram-stain brightfield preparation compares the two size classes, huge and standard, of Gram-negative spirilla. (Bar $-5 \mu \text{m.}$) (k) Standard-sized spirillum Gram stain. The lighter spots are probably sulfur globules. (Bar $-5 \mu \text{m.}$)

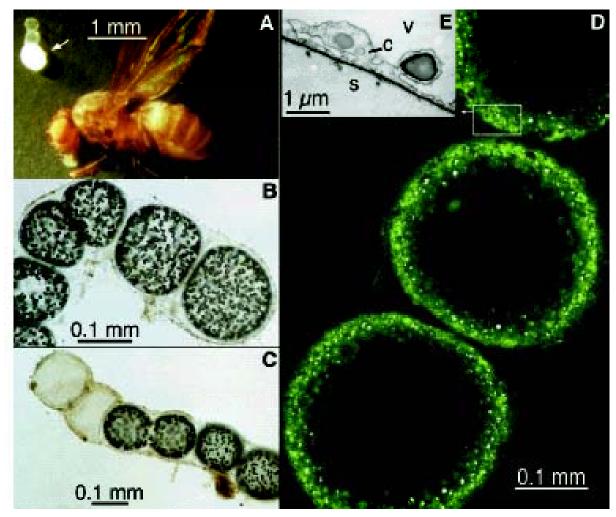
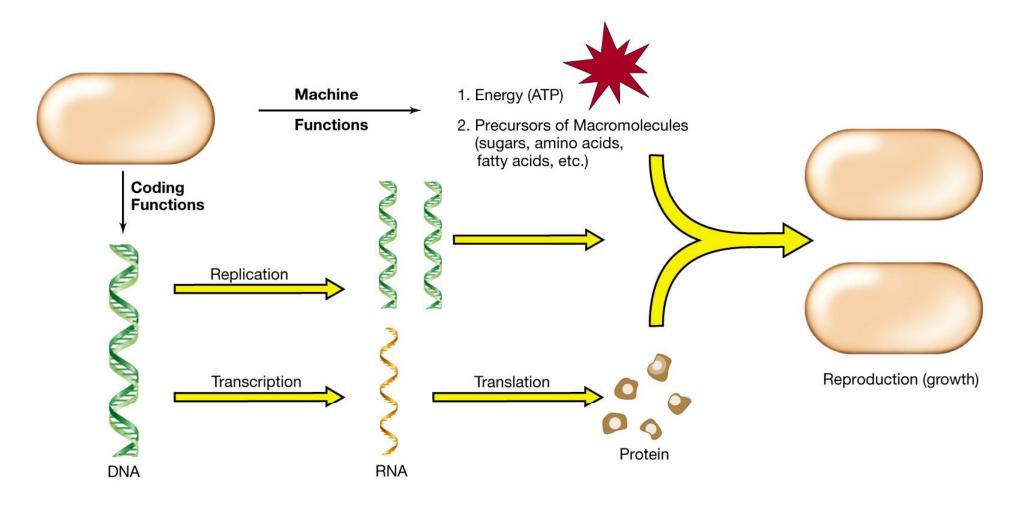


Fig. 1. Thiomargarita namibiensis. (A) The white arrow points to a single cell of Thiomargarita, 0.5 mm wide, which shines white because of internal sulfur inclusions. Above there is an empty part of the sheath, where the two neighboring cells have died. The cell was photographed next to a fruit fly (Drosophila viriles) of 3 mm length to give a sense of its size. (B) A typical chain of Thiomargarita as it appears under light microscopy. (C) At the left end of the chain there are two empty mucus sheaths, while in the middle a Thiomargarita cell is dividing. (D) Confocal laser scanning micrograph showing cytoplasm stained green with fluorescein isothiocyanate and the scattered light of sulfur globules (white). Most of the cells appear hollow because of the large central vacuole. (E) Transmission electron micrograph of the cell wall [enlarged area in (D)] showing the thin layer of cytoplasm (C), the vacuole (V), and the sheath (S).

Thiomargarita namibiensis

Up to 500 µm wide

The machine/coding functions of the cell



Central Dogma

Chemical features of a "typical" bacterial cell (E. coli)

Table 3.1 Molecular composition of a bacterial cell, Escherichia coli, during balanced exponential growth.^a

Component	Percentage of total weight ^b	Approximate number of molecules/cell	Number of different kinds
Water	70%	20,000,000,000	1
Proteins	16%	2,400,000	2,000°
RNA: rRNA, tRNA, and other small regulatory RNA (sRNA)		250,000	200
mRNA	0.7%	4.000	2,000°
Lipids: phospholipids (membrane)	3%	25,000,000	50 Mos
lipopolysaccharide (outer membrane)	1%	1,400,000	1 rRN
DNA	1%	2 ^d	
Metabolites and biosynthetic precursors	1.3%	50,000,000	1,000
Peptidoglycan (murein sacculus)	0.8%	1	
Inorganic ions	0.1%	250,000,000	20
Polyamines (mainly putrescine and spermine)	0.1%	6,700,000	2

^aValues shown are for a hypothetical "average" cell cultured with aeration in glucose medium with minimal salts at 37°C.

^bThe total weight of the cell (including water) is about 10⁻¹² gram (g), or 1 picogram (pg).

^cThe number of kinds of mRNA and of proteins is difficult to estimate because some genes are transcribed at extremely low levels and because RNA and proteins include kinds that are rapidly degraded.

In rapidly growing cells, cell fission typically lags approximately one generation behind DNA replication; hence, two identical DNA copies per cell.

Source: Modified from Neidhardt, F., and H. E. Umbarger. 1996. Chemical composition of *Escherichia coli*, p. 14. In *Escherichia coli* and *Salmonella*: Cellular and Molecular Biology, 2nd ed. ASM Press, Washington, DC.

Table 3-1 Microbiology: An Evolving Science © 2009 W. W. Norton & Company, Inc.

Take Home Message: Proteins are #1 by weight Lipids are #1 by number Peptidoglycan is 1 jumbo molecule RNA is mostly ribosomes DNA is also a huge polymer

Macromolecules in a typical bacterial cell:

Not just "soup" – highly ordered cytoplasm

Contents (e.g., proteins) will vary, depending on conditions.

Z

mRNA

Flagellar

motor

Flagellum

DNA-binding protein

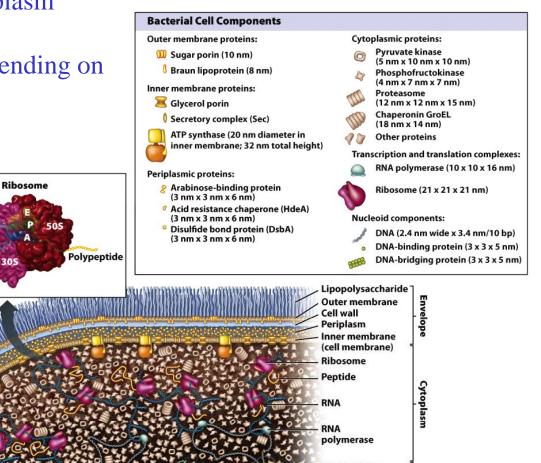
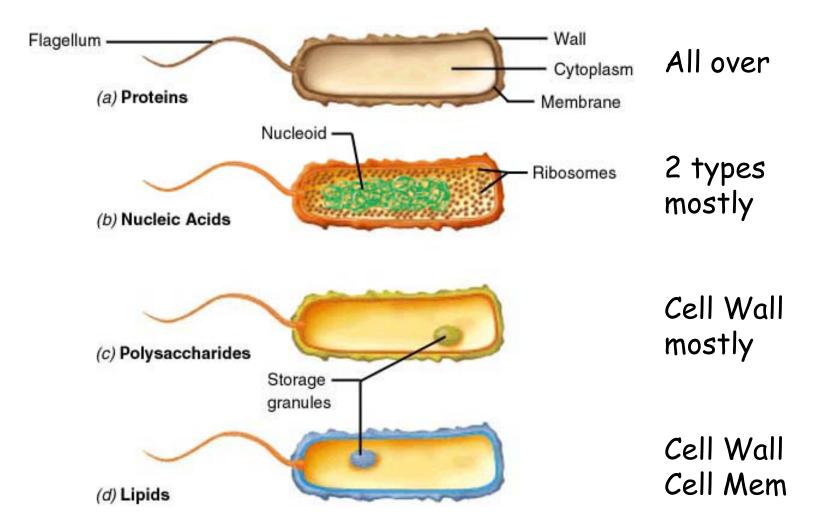




Figure 3.1 part 2 Microbiology: An Evolving Science © 2009 W.W. Norton & Company, Inc.

Locations of macromolecules in the cell



Comparing Bacteria, Archaea and Eucarya

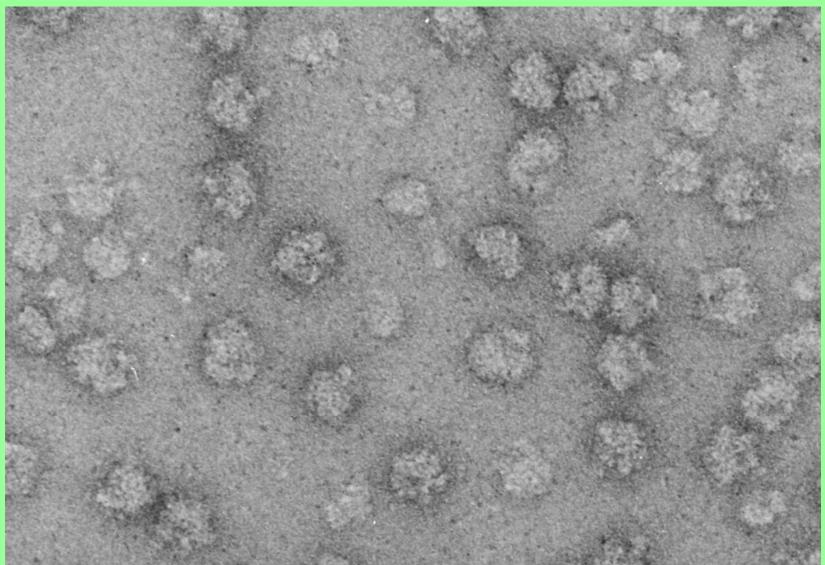
Classification of microbial cellular features: Invariant (or common to all)

Ribosomes: Sites for protein synthesis – aka the grand translators.

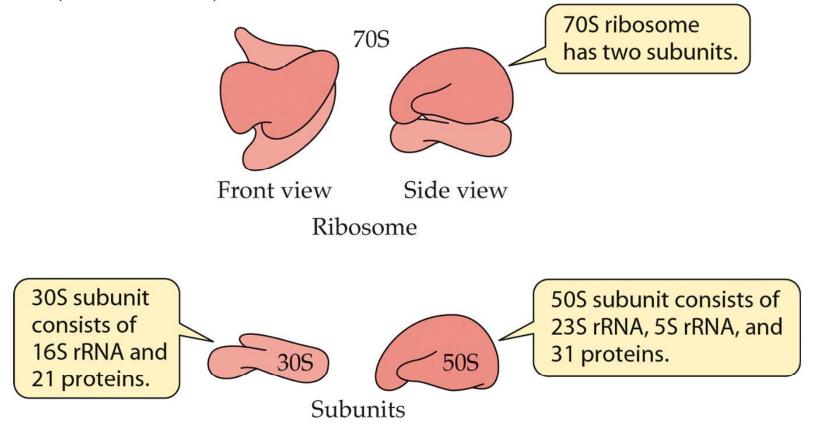
Cell Membranes: The barrier between order and chaos.

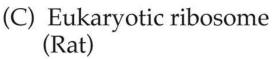
Nucleoid Region: Curator of the Information.

Ribosome structure



(B) Prokaryotic ribosome (*Escherichia coli*)





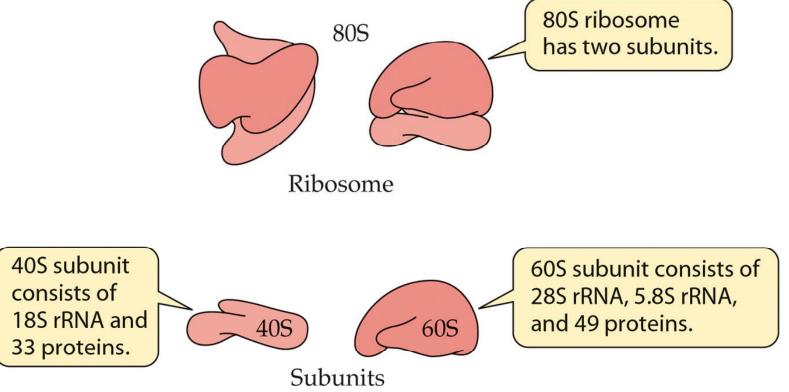


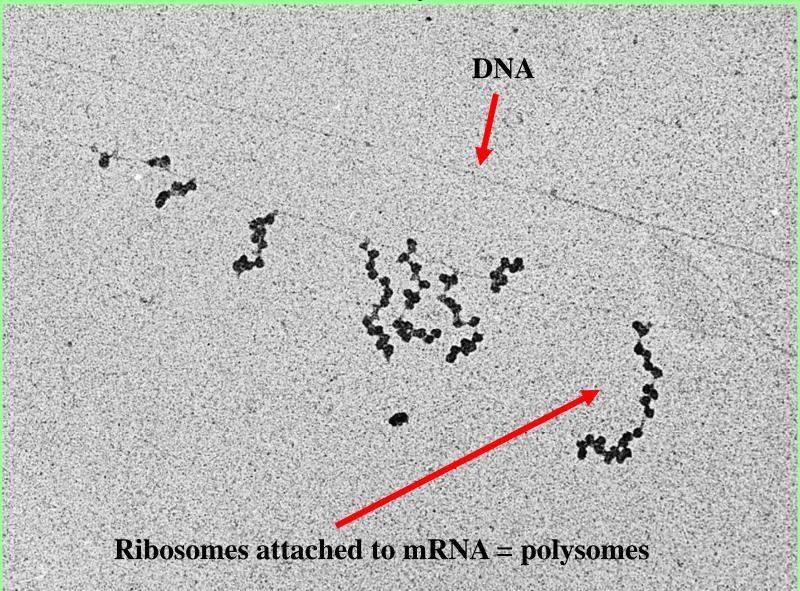
Table 7.6Ribosome structure^a

Property	Prokaryote	Eukaryote
Overall size	70S	80S Most Complex
Small subunit	30S	40S
Number of proteins	~21	~ 30
RNA size (number of bases)	16S (1500)	18S (2300)
Large subunit	50S	60S
Number of proteins	~ 34	~ 50
RNA size (number of bases)	23S (2900)	28S (4200)
	5S (120)	5.8S (160)
		5S (120)

^{*a*} Ribosomes of mitochondria and chloroplasts of eukaryotes are similar to prokaryotic ribosomes (*Construction* 14.4).

S= Svedberg; a sedimentation coefficient that is NOT ADDITIVE!!!

Protein synthesis



Comparing Bacteria, Archaea and Eucarya

Classification of microbial cellular features: Invariant (or common to all)

Ribosomes: Sites for protein synthesis – aka the grand translators.

Cell Membranes: The barrier between order and chaos.

Nucleoid Region: Curator of the Information.

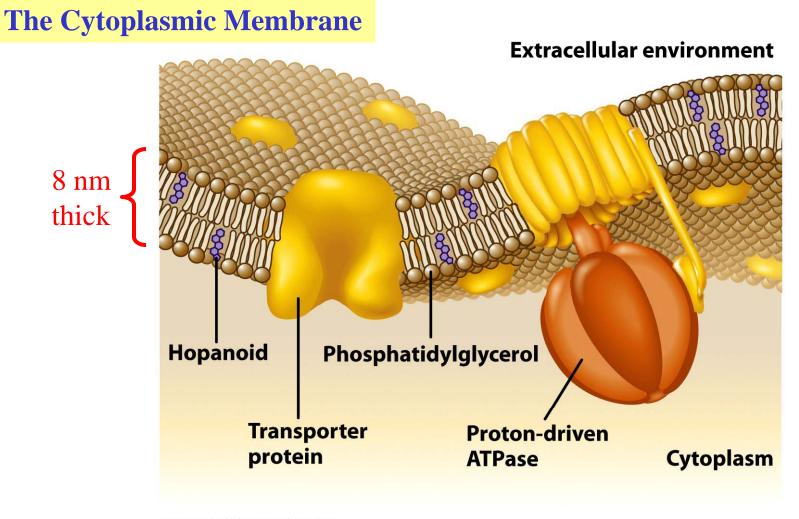
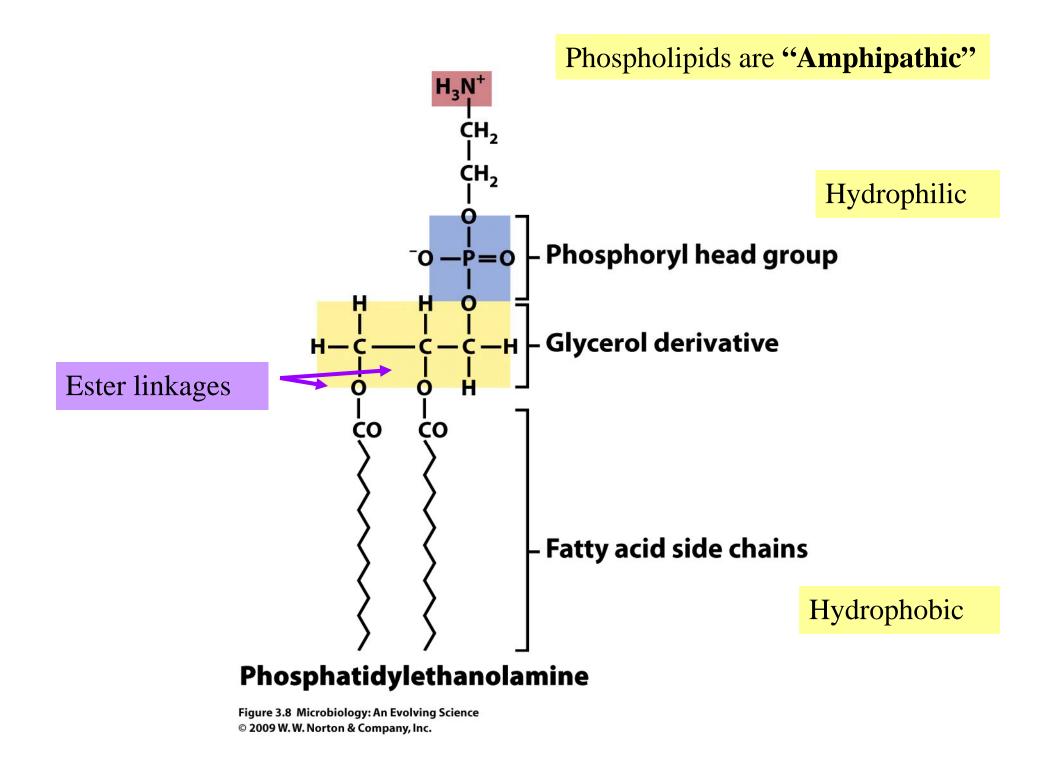
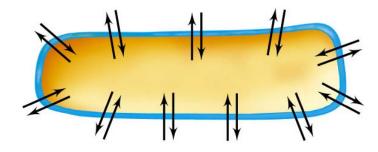


Figure 3.7 Microbiology: An Evolving Science © 2009 W. W. Norton & Company, Inc.

Membrane has similar viscosity as oil: Fluid Mosaic Model

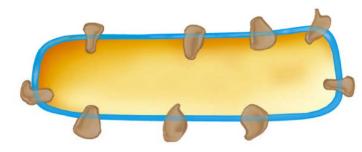
Stabilized by H bonds, hydrophobic interactions, and by Mg^{++} and Ca^{++} binding to phosphate heads



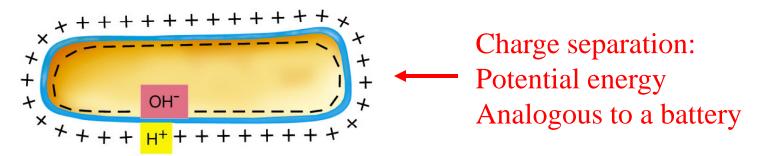


Functions of the cytoplasmic membrane

Permeability Barrier — Prevents leakage and functions as a gateway for transport of nutrients into and out of the cell



Protein Anchor — Site of many proteins involved in transport, bioenergetics, and chemotaxis



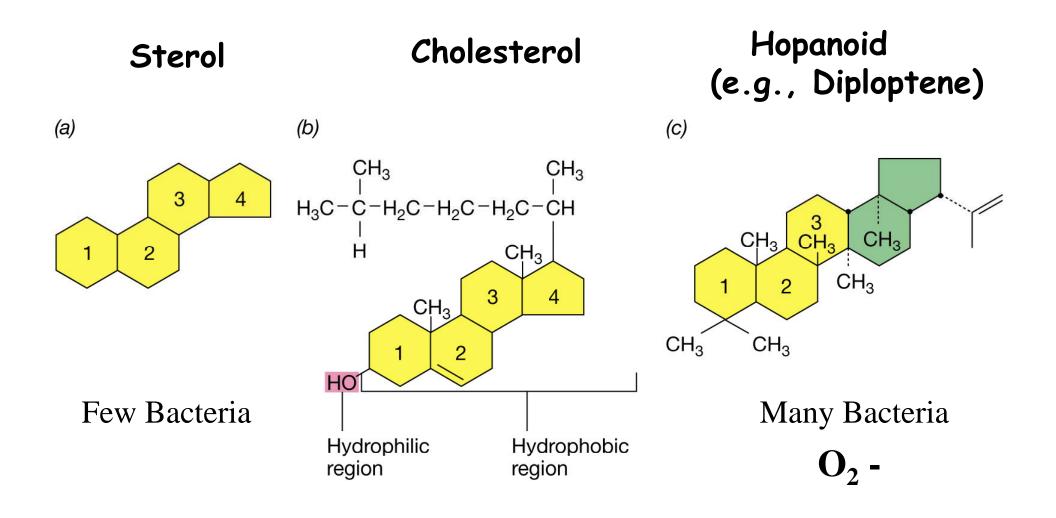
Energy Conservation — Site of generation and use of the proton motive force

Comparative permeability of membranes to various molecules

Rate of permeability ^a		
100	Free diffusion of water (passive transport)	
0.1 0.001	assisted by aquaporins	
0.001		
	Active or passive transport (depends on conditions)	
0.00000001	(depends on conditions), but proteins aid movement	
	100 0.1 0.001 0.0001 0.000001 0.0000001	

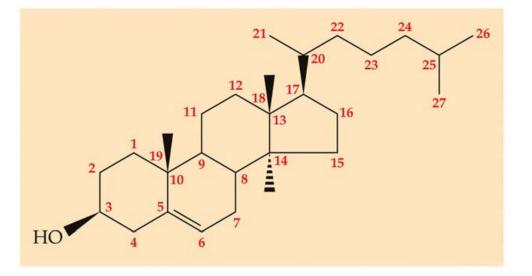
Table 4.2

^{*a*} Relative scale—permeability with respect to permeability of water given as 100. Permeability of the membrane to water may be affected by aquaporins (see text).

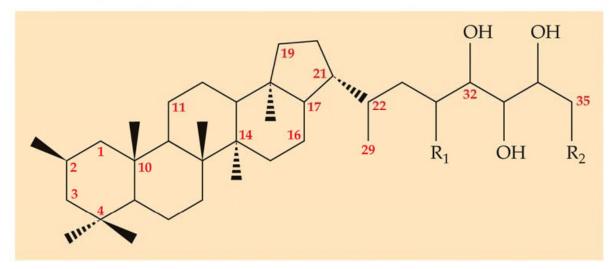


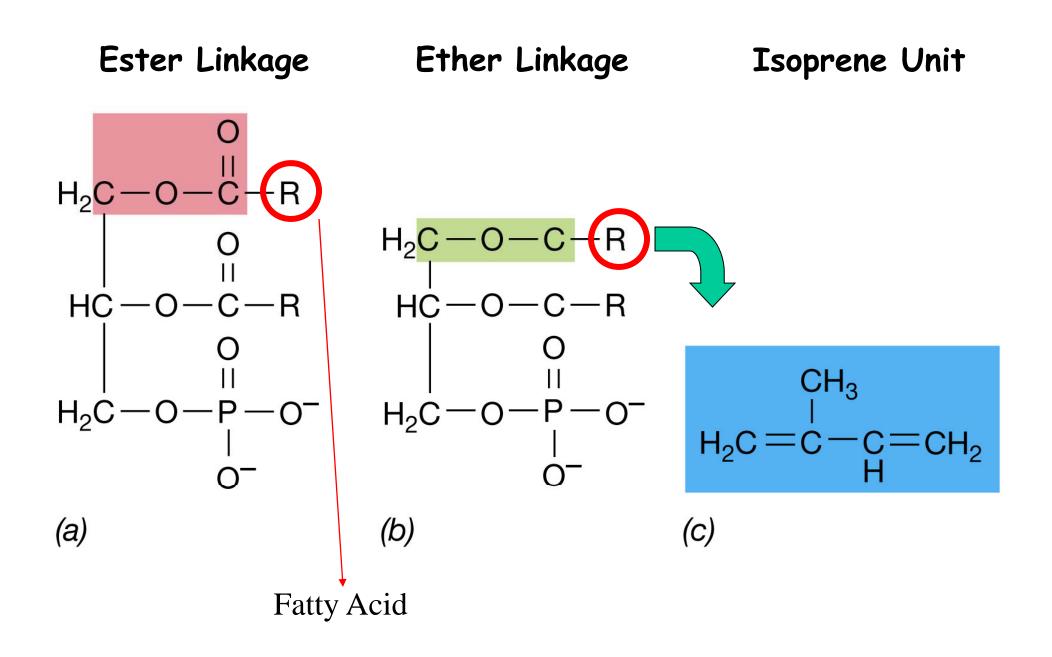
All rigid planar molecules

(A) Cholesterol

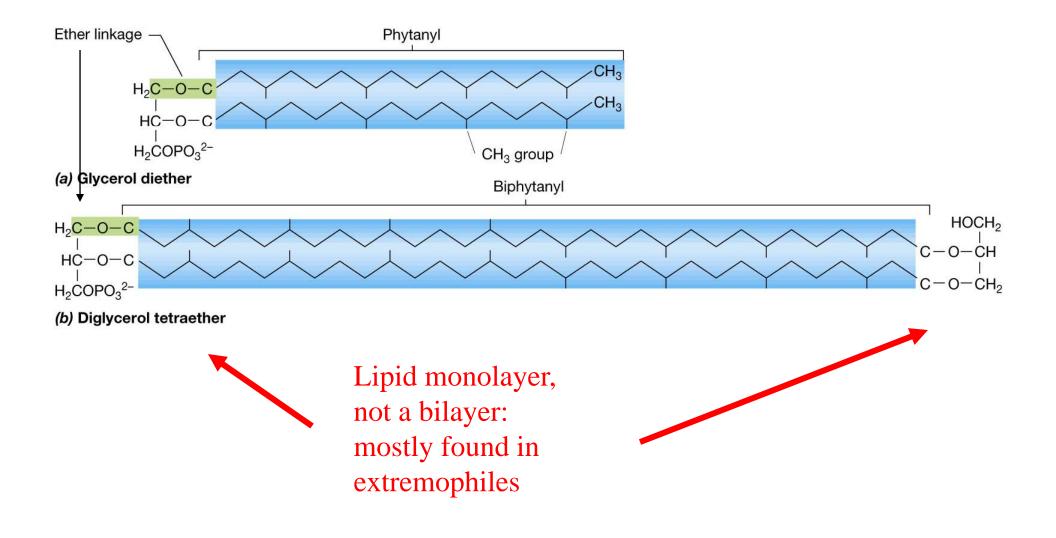


(B) A hopanoid from a cyanobacterium

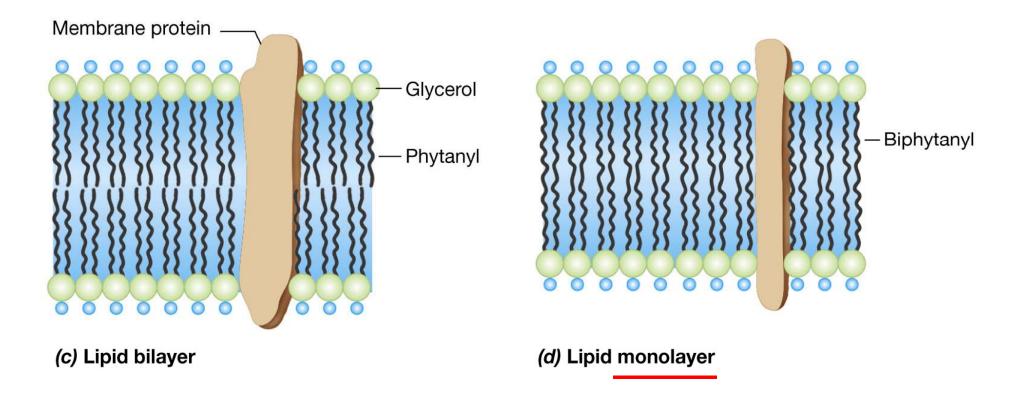




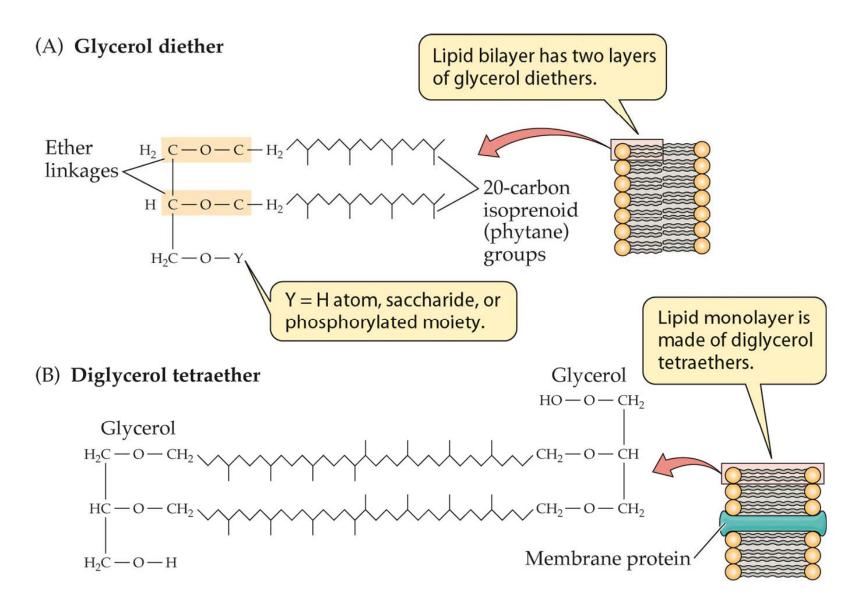
Major lipids of *Archaea* and the structure of archaeal membranes



Major lipids of *Archaea* and the structure of archaeal membranes



Archaeal cell membrane structure



Passive Diffusion:

-small, uncharged molecules (O₂, CO₂, H₂O)

-weak acids and bases in protonated form (more hydrophobic)

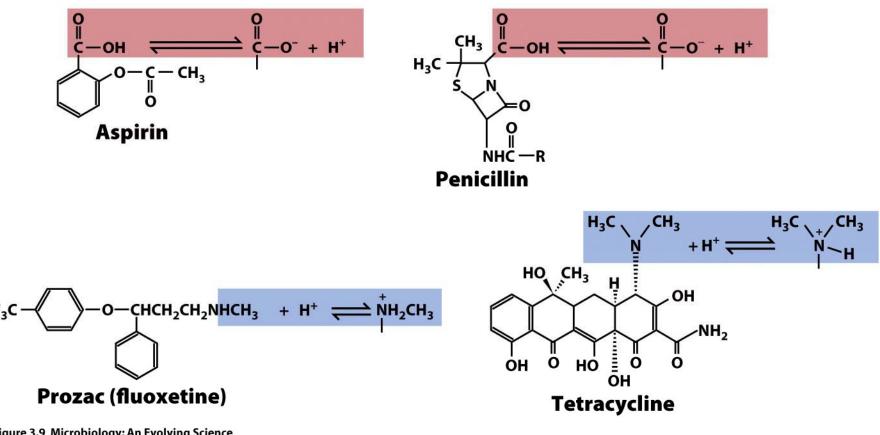


Figure 3.9 Microbiology: An Evolving Science © 2009 W. W. Norton & Company, Inc.

Facilitated Diffusion (a type of passive tranport): Powered by solute's own concentration gradient

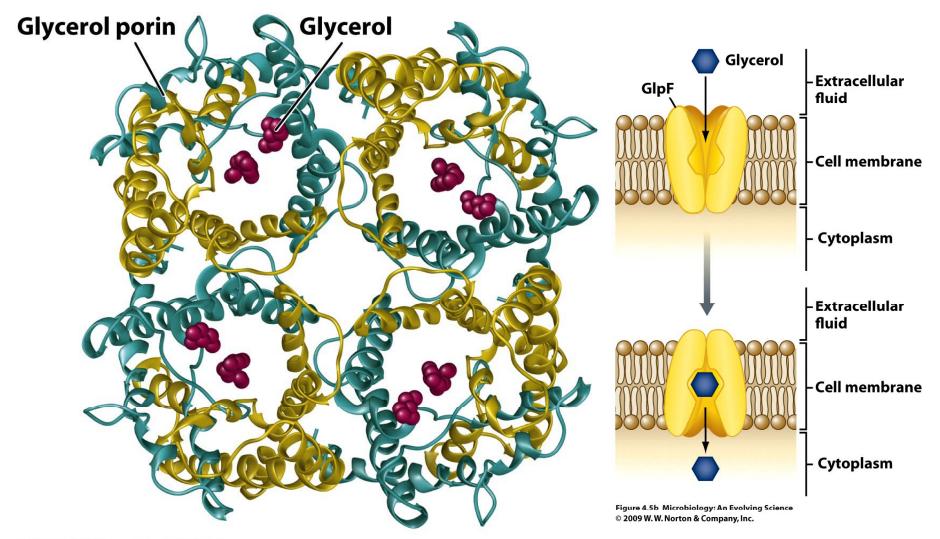


Figure 4.5a Microbiology: An Evolving Science © 2009 W. W. Norton & Company, Inc.

Aquaporins, glycerol transporters

But how do you get glucose into the cell? The overall strategy for feeding in Bacteria and Archaea:

Suppose you wished to use bacteria in a landfill to break down paper. Paper consists of a polymer, cellulose. It is a readily metabolizable macromolecule - it is a good *carbon and energy source* for bacteria

<u>Problem</u>: Microbes do not ingest – no phagocytosis, no pinocytosis – everything must be solubilized before transport into the cell. How do you get the cellulose into the cell?

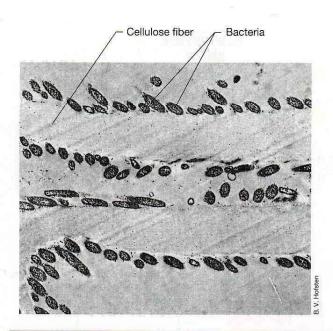
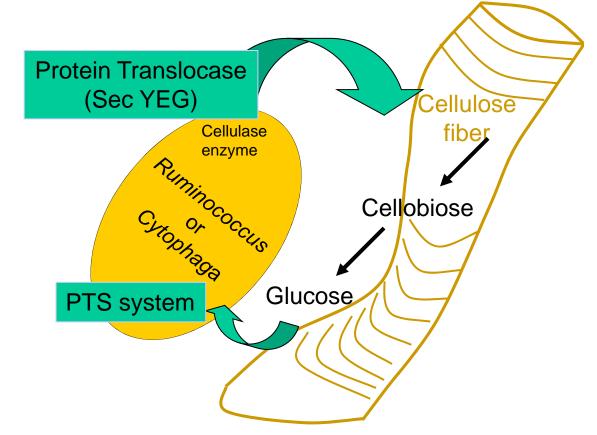
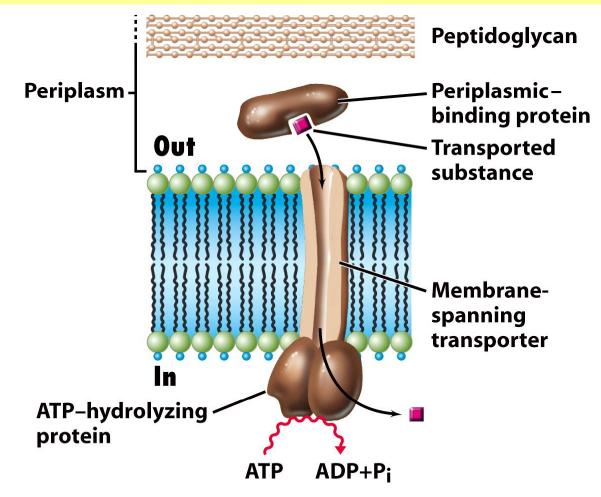


Figure 17.61 Transmission electron micrograph showing attachment of cellulose-digesting bacteria, *Sporocytophaga myxococcoides*, to cellulose fibers. Cells are about 0.5 µm in diameter.



Primary Transport (a type of **Active Transport**):

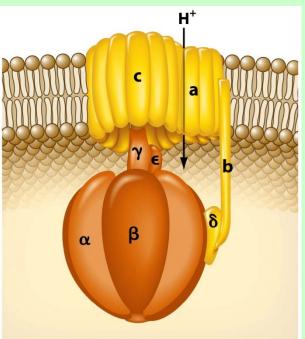
- \rightarrow ATP hydrolysis provides energy for transfer
- \rightarrow ABC transporters = <u>ATP-B</u>inding <u>Cassette</u>
- 1. Periplasmic binding proteins "find" low-concentration solutes ($\geq 10^{-6}$ M)
- 2. Binding protein docks on membrane-spanning domain
- 3. Conformational change and ATP hydrolysis drive transport



Proton pumps (e.g. cytochrome oxidase) push protons out of cell; the electron transport chain is **anchored in membrane**.

Energy conservation: proton-motive force (PMF) is generated from protons.

- Osmotic force tries to push protons back into cell
- Electrical force tries to push protons back into cell



PMF is used to create ATP via the enzyme ATP synthase

Comparing Bacteria, Archaea and Eucarya

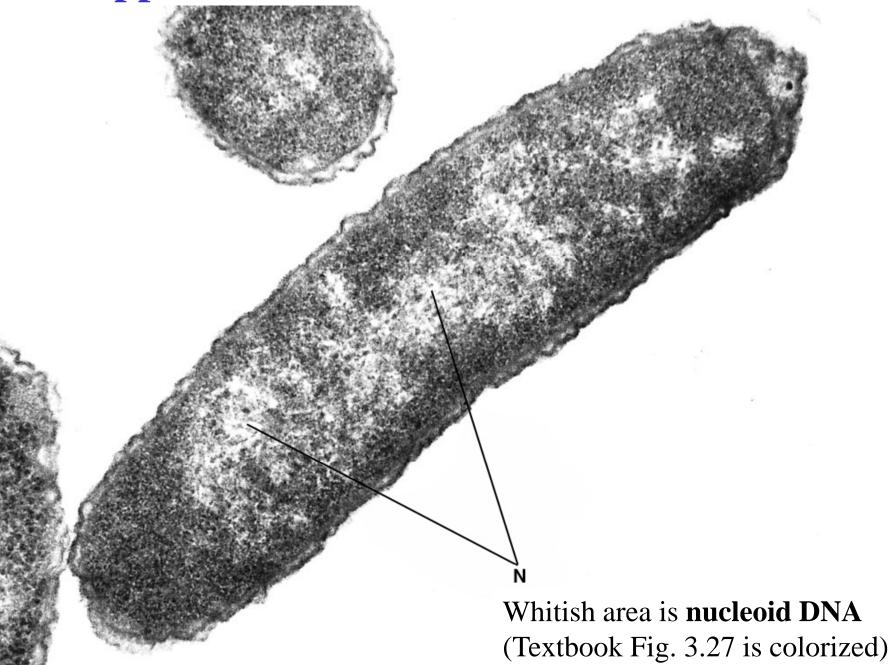
Classification of microbial cellular features: Invariant (or common to all)

Ribosomes: Sites for protein synthesis – aka the grand translators.

Cell Membranes: The barrier between order and chaos.

Nucleoid Region: Curator of the Information.

Appearance of nucleoid via TEM



DNA strands released from cell



Bacterial & Archaeal DNA

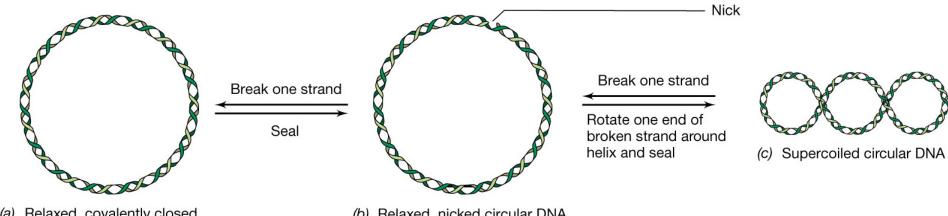
Statistics:

Chromosomes: ~1; bears essential genes Plasmids: 0 to hundreds; helpful genes Circumference: ~1 mm



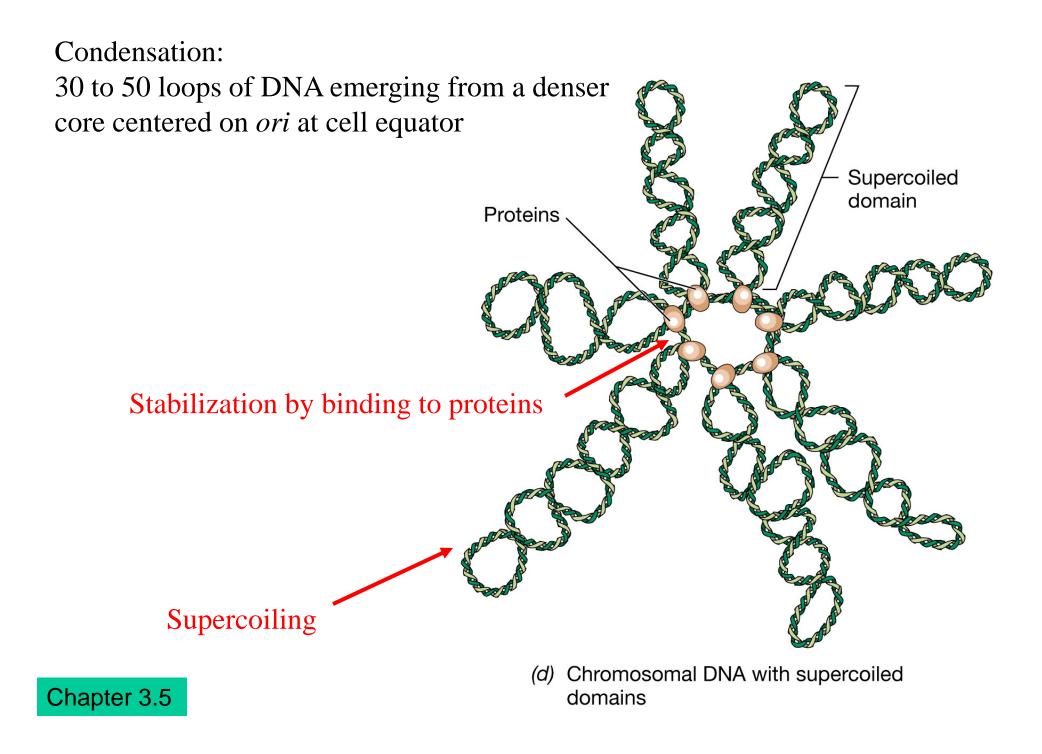
Enigma:

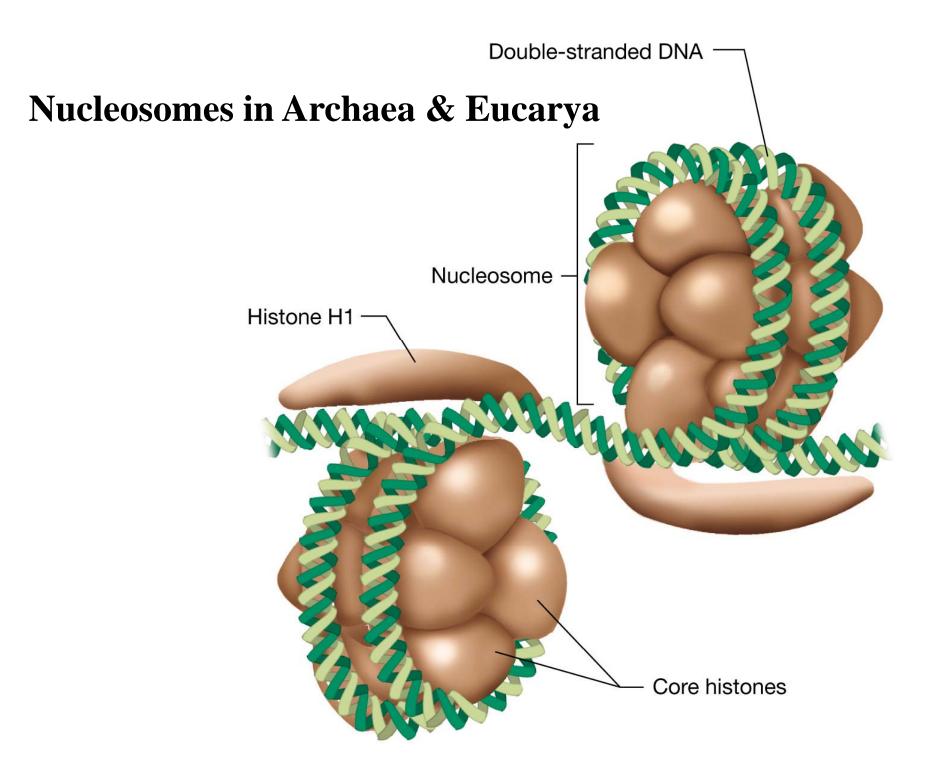
How to fit 1 mm long chromosome into a 1 µm wide cell? Condensation: 30 to 50 loops of DNA emerging from a denser core Supercoiling: tight twisting Organization: wrapped around histone-like proteins (in Bacteria) or histones (in Archaea)



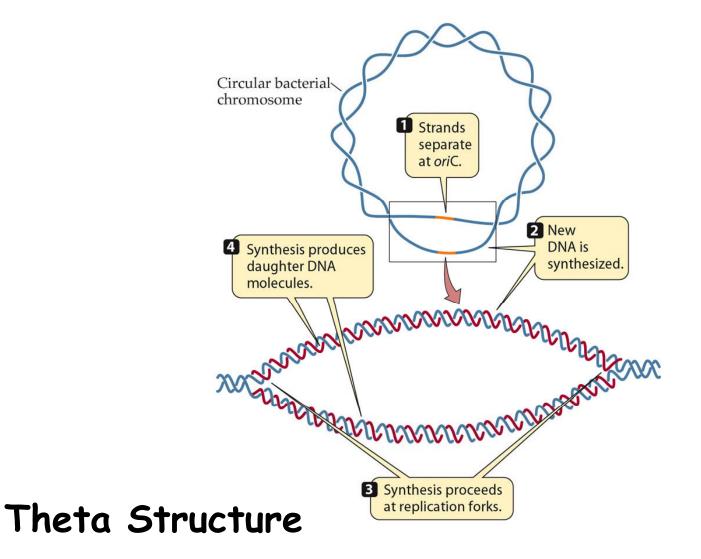
(a) Relaxed, covalently closed circular DNA

(b) Relaxed, nicked circular DNA





Overview of DNA replication

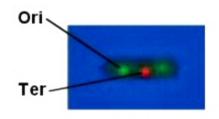


See Chapter 7.3, 7.5 - 7.7

Overview of DNA replication

Fluorescence microscopy: *E. coli* cells with fluorophores labeling Ori and Ter

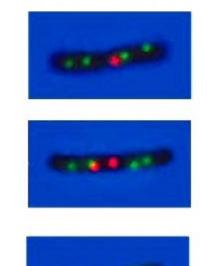
Lau et al (2003) Mol. Micro. 49:731

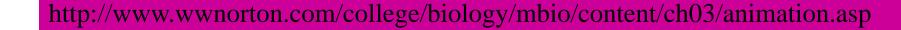


40 minutes to replicate *E. coli* chromosome.

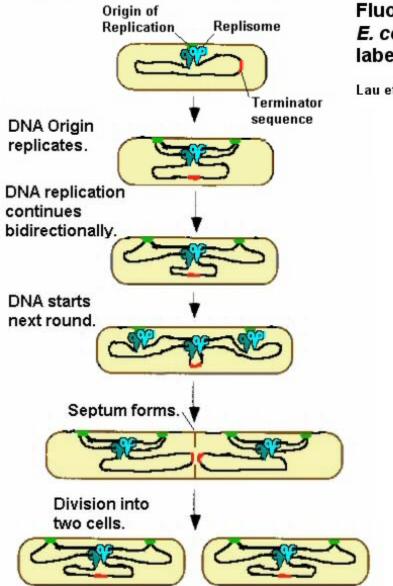
20 minutes for cell division.

How???



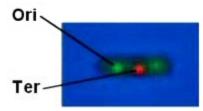


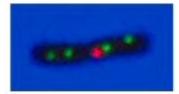
Overview of DNA replication

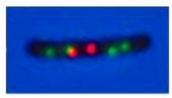


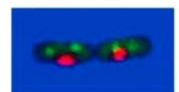
Fluorescence microscopy: *E. coli* cells with fluorophores labeling Ori and Ter

Lau et al (2003) Mol. Micro. 49:731

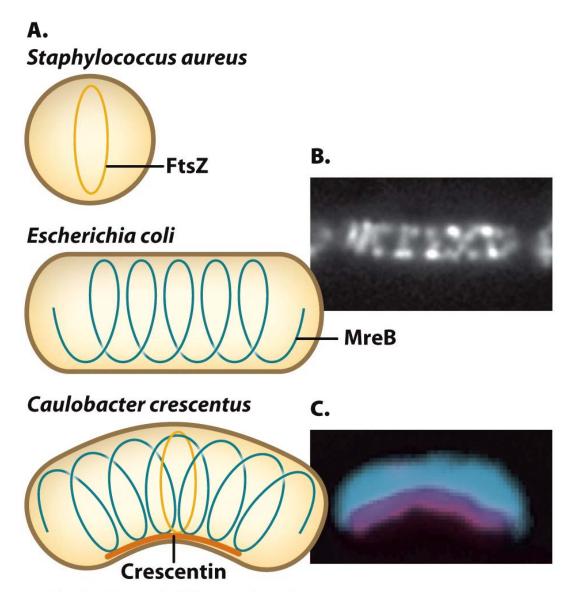








Bacteria have cytoskeletons!



Special Topic 3.2 figure 2 Microbiology: An Evolving Science © 2009 W.W. Norton & Company, Inc. Photos: N. Ausmees, et al. 2003

Bacteria divide by binary fission. FtsZ marks the spot.

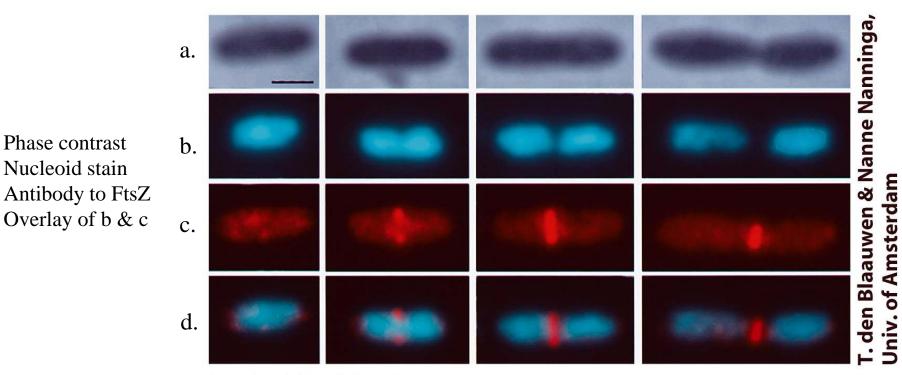


Figure 6-2b Brock Biology of Microorganisms 11/e © 2006 Pearson Prentice Hall, Inc.

FtsZ is a structural analog (ancestor?) of eukaryotic tubulin

 \rightarrow forms Z-ring at future site of cytokinesis

Phase contrast

Nucleoid stain

a.

b.

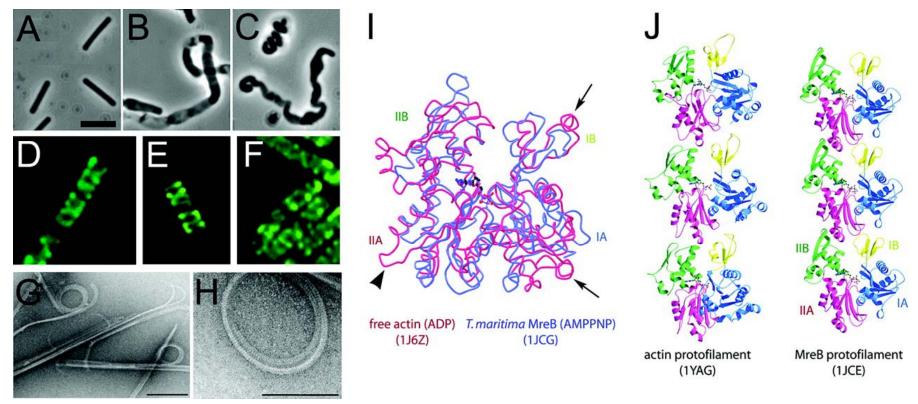
c.

d.

 \rightarrow only 17% amino acid identity to tubulin but similar 3D structures and assembly properties

MreB is a homolog (ancestor?) of actin

-cell shape determinant-present in rod- and spiral-shaped cells but absent from cocci-only 15% amino acid identity but similar 3D structure



A-C: WT and mreB mutants of *B. subtilis* (note cell shapes)

D-F: Helical filaments formed by MreB-like proteins in B. subtilis

G&H: MreB filaments

I &J: Actin and MreB structures overlaid. Only 15% amino acid identity but similar 3D structure.



Gemmata obscuriglobus

Membrane encompassed nucleoid