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**Microbial Taxonomy**

Traditional taxonomy or the classification through **identification** and **nomenclature** of microbes, both "prokaryote" and eukaryote, has been in a mess – we were stuck with it for traditional reasons.

A "natural" taxonomy would be based on evolutionary relatedness: Thus, organisms in same "genus" (a collection of "species") would have similar properties in a fundamental sense.

A natural taxonomy of macrobes has long been possible: Large organisms have many easily distinguished features (e.g., body-plans and developmental processes, that can be used to describe hierarchies of relatedness).

Microbes usually have few distinguishing properties that relate them, so a hierarchical taxonomy mainly has not been possible.

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Recent advances in **molecular phylogeny** have changed this picture. We now have a relatively quantitative way to view **biodiversity**, in the context of phylogenetic maps or evolutionary trees.

Slowly evolving molecules (e.g. rRNA) used for large-scale structure; "fast- clock" molecules for fine-structure.

The literature language (e.g. "species") and formal nomenclature, however, remain solidly rooted in the tradition of Linnaeus at this time. (You have to call them something!)

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**Table 11.4 Some phenotypic characteristics of taxonomic value**

Major category	Components
I. Morphology	Shape; size; Gram reaction; arrangement of flagella, if present
II. Motility	Motile by flagella; motile by gliding; motile by gas vesicles; nonmotile
III. Nutrition and Physiology	Mechanism of energy conservation (phototroph, chemosynotroph, chemolithotroph); relationship to oxygen; temperature, pH, and salt requirements/tolerances; ability to use various carbon, nitrogen, and sulfur sources; growth factor requirements
IV. Other factors	Pigments; cell inclusions, or surface layers; pathogenicity; antibiotic sensitivity

**Enterotube™ reactions**

Uninoculated Enterotube™  
After 24h growth of *Salmonella enterica*

Species	+ Glucose acid	Lipase	Urease	Indole	Lactose	α-Glycerol phosphate	α-Ketoglutarate	Starch
<i>Salmonella enterica</i>	0.99	0.98	0.97	0.95	0.91	0.91	0.95	0.96
<i>Proteus mirabilis</i>	0.99	0.91	0.99	0.98	0.91	0.93	0.91	0.91
<i>Shigella sonnei</i>	0.99	0.91	0.95	0.91	0.91	0.95	0.99	0.91
<i>Enterobacter aerogenes</i>	0.99	0.98	0.98	0.91	0.95	0.95	0.99	0.91
<i>Pseudomonas fluorescens</i>	0.99	0.98	0.91	0.91	0.98	0.98	0.99	0.91

Substrates: Lactose, Arabinose, Sorbitol, Voges-Proskauer, Indole/phenylalanine, Urea, Citrate

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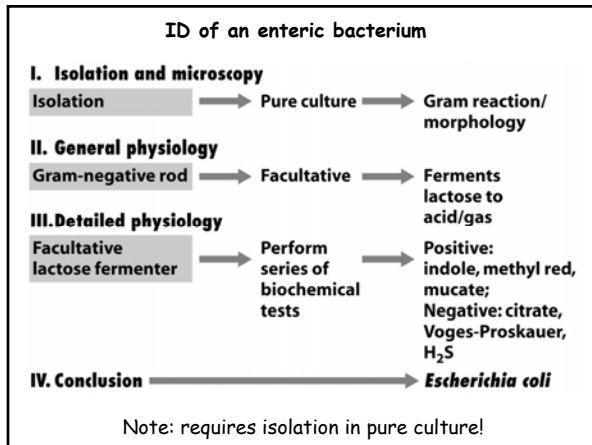
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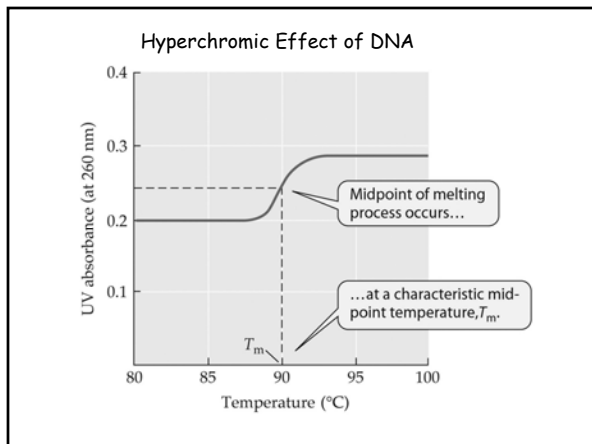
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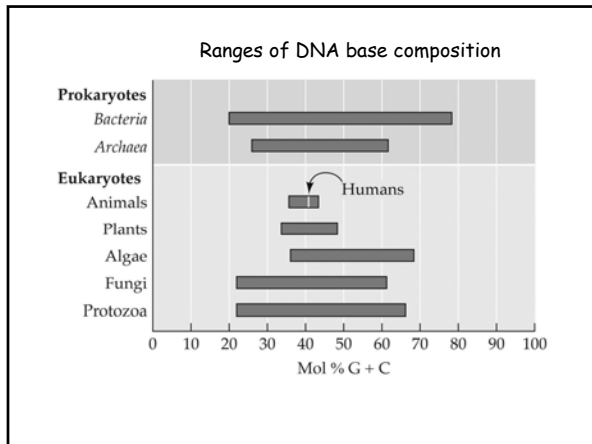
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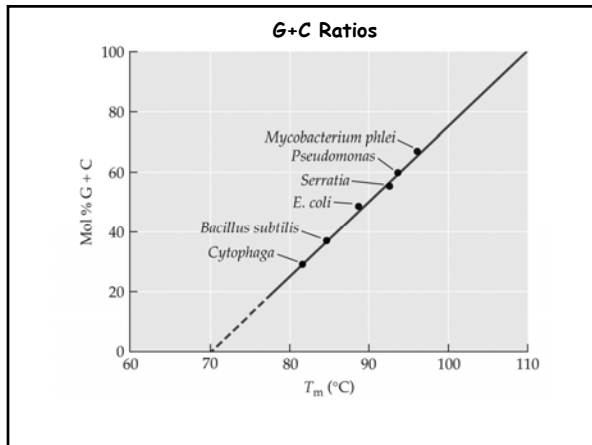
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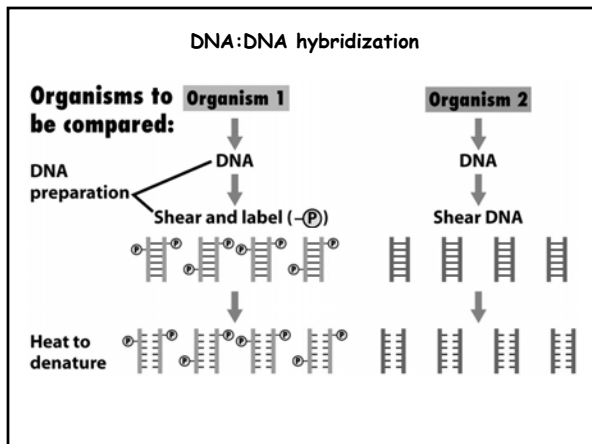
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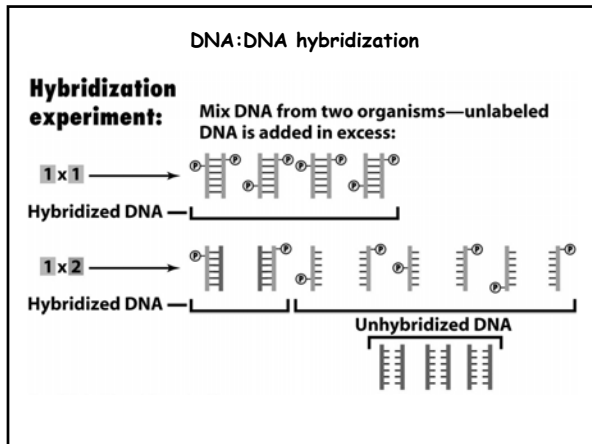
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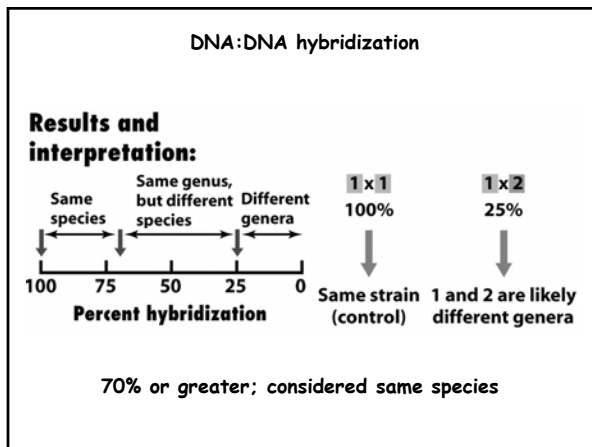
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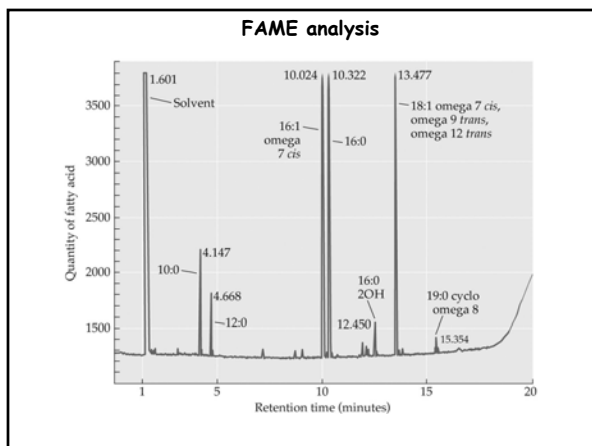
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**Classes of Fatty Acids in Bacteria**

Class/Example	Structure of example
I. <b>Saturated:</b> tetradecanoic acid	$\text{HO}-\text{C}(=\text{O})-(\text{CH}_2)_{12}-\text{CH}_3$
II. <b>Unsaturated:</b> <i>omega</i> -7- <i>cis</i> hexadecanoic acid	$\text{HO}-\text{C}(=\text{O})-(\text{CH}_2)_6-\text{C}(\text{H})=\text{C}(\text{H})-(\text{CH}_2)_6-\text{CH}_3$
III. <b>Cyclopropane:</b> <i>cis</i> 7, 8 methylene hexadecanoic acid	$\text{HO}-\text{C}(=\text{O})-(\text{CH}_2)_7-\text{C}(\text{H})_2-\text{C}(\text{H})_2-(\text{CH}_2)_5-\text{CH}_3$
IV. <b>Branched:</b> 13-methyltetradecanoic acid	$\text{HO}-\text{C}(=\text{O})-(\text{CH}_2)_{10}-\text{C}(\text{H})(\text{CH}_3)-\text{CH}_3$
V. <b>Hydroxy:</b> 3-hydroxytetradecanoic acid	$\text{HO}-\text{C}(=\text{O})-\text{CH}_2-\text{C}(\text{H})(\text{OH})-(\text{CH}_2)_{10}-\text{CH}_3$

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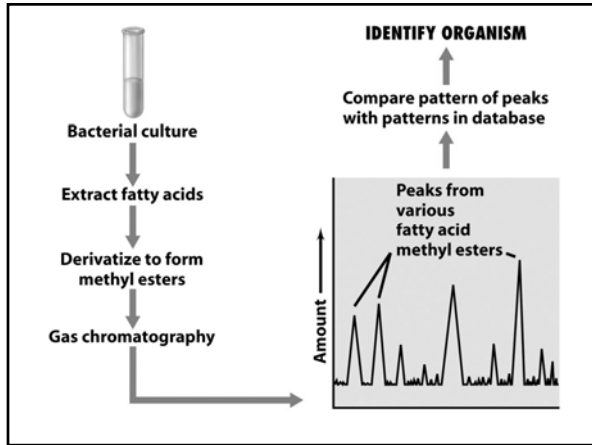
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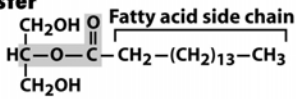
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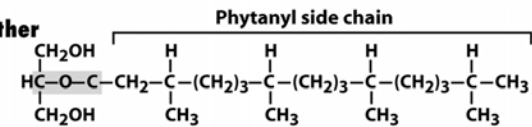
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**Ester**



**Bacteria, Eukarya**

**Ether**



**Archaea**

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**Table 17.1** Hierarchical classification of the bacterium *Spirochaeta plicatilis*

Taxon	Name
Domain	<i>Bacteria</i>
Phylum	<i>Spirochaetes</i> (vernacular name: spirochetes)
Class	<i>Spirochaetes</i>
Order	<i>Spirochaetales</i>
Family	<i>Spirochaetaceae</i>
Genus	<i>Spirochaeta</i>
Species	<i>plicatilis</i>

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**Table 17.2** Taxonomic hierarchy of classification.

Taxon rank	A long-studied taxon	A less-studied taxon	An uncultivated environmental sample
Domain	Bacteria	Archaea	Bacteria
Division	Actinobacteria	Euryarchaeota	Proteobacteria
(phylum)	Filamentous gram-positive	Methanogens and halophiles	Purple bacteria and relatives; gram-negative
Class	Actinobacteria	Methanococci	Alpha Proteobacteria
	High GC gram-positive	Methanogens	Gram-negative bacteria
Subclass	Actinobacteridae		
Order	Actinomycetales	Methanococcales	Rickettsiales
	Filamentous; acid-fast stain	Methanogenic cocci	Includes intracellular bacteria
Suborder	Streptomycinae		
Family	Streptomycetaceae	Methanocaldococcaceae	SAR11 cluster
	Filamentous; hyphae produce spores	Thermophilic methanogens	Nonculturable planktonic marine bacteria
Genus	<i>Streptomyces</i>	<i>Methanocaldococcus</i>	<i>Pelagibacter</i>
Species	<i>S. coelicolor</i>	<i>M. jamaecum</i>	<i>P. ubique</i>
(date first described)	(1908)	(1984)	(2002)

Table 17.2 Microbiology: An Evolving Science  
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**Table 11.6** Taxonomic ranks and numbers of known prokaryotic species<sup>a</sup>

Rank	Bacteria	Archaea	Total
Domains	1	1	2
Phyla	25	4 <sup>a</sup>	29
Classes	34	9	43
Orders	78	13	91
Families	230	23	243
Genera	1227	79	1306
Species	6740	289	7029

<sup>a</sup>Numbers represent validly named genera and species of *Bacteria* and *Archaea* as of 2005. The phyla category for *Archaea* includes the Korarchaeota and the Nanoarchaeota, not yet officially recognized phyla.

Source: Garrity, G.M., Libum, T.G., and Bell, J.A. 2005. *Bergey's Manual of Systematic Bacteriology*, 2d ed., Vol. 2, part A, pp159–220. Springer-Verlag, New York.

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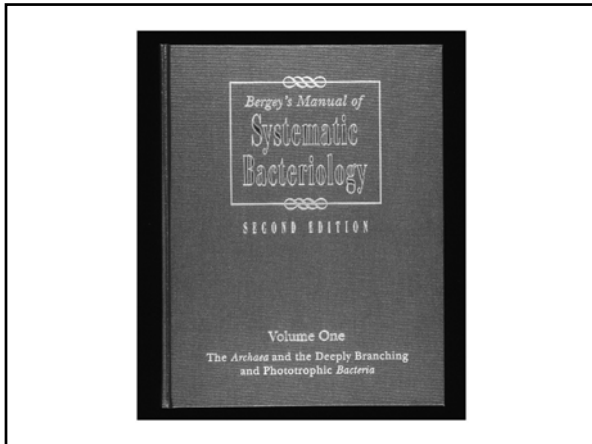
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**Taxonomy Summary**

Classical physiological descriptions of microbes constitute a taxonomy, but do not provide relationships (except as might be inferred subjectively).

Methods such as G+C ratios, FAME, DNA-DNA hybridization establish relationships, but only if close, i.e., they are not sufficiently general to be broadly applicable.

All these methods require pure-cultivation of organisms for characterization, but we can't cultivate much of what is out there.

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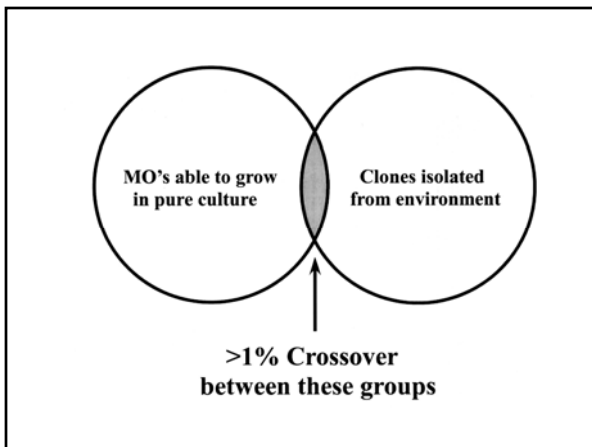
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## Importance of a Molecular Biological Approach

- **Traditional culturing** techniques isolate ~1% of the total bacteria in marine ecosystems, thereby severely underestimating diversity and community structure.
- Because nutrient-rich **culture media** have been historically used during enrichment procedures, bacteria which may be dominant in natural communities are selected against in favor of copiotrophic (weedy) bacteria.
- **SSU rRNAs** and their respective genes are excellent descriptors of microbial taxa based on phylogeny.

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Stanier *et al.*, 1976:

*... the general course of evolution [for bacteria] will probably never be known, and there is simply not enough objective evidence to base their classification on phylogenetic grounds...*

*For these and other reasons, most modern taxonomists have explicitly abandoned the phylogenetic approach.*

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## Regarding Molecular Phylogeny

### The Root of the Problem:

Unlike zoology and botany, microbiology developed without the knowledge of phylogenetic relationships among the organisms studied.

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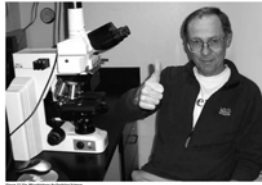


## Molecular Phylogeny



← Woese (1977): Applied rRNA concept to redefine microbial systematics (microbial genealogy).

→ Pace (1984): Applied rRNA concept to microbial ecology (census without culturing).



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## Why ribosomal RNAs?

- Found among all living organisms (for 3.8 of the last 4.5 billion years). Integral part of protein synthesis machinery.
- Cell component analyses provide culture-independent means of investigating questions in microbial ecology (lack of morphology).
- rRNAs offer a type of sequence information that makes them excellent descriptors of an organism's evolutionary history.
- No detectable horizontal gene transfer, especially important for the bacteria and archaea.
- Large and growing database; RDP contains  $\sim 1.3 \times 10^6$  SSU rRNAs.

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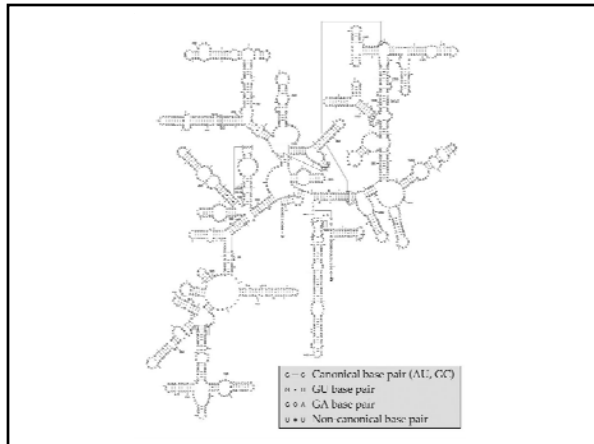
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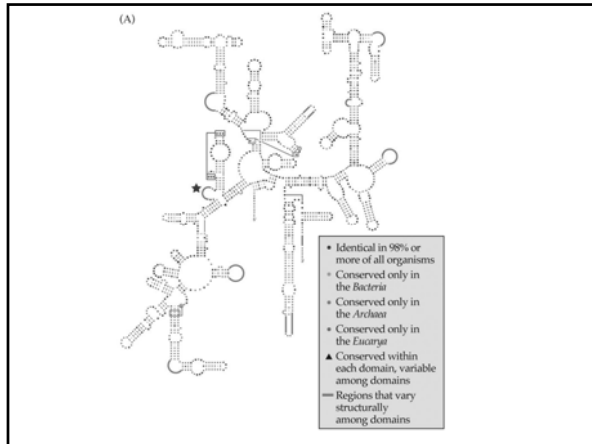
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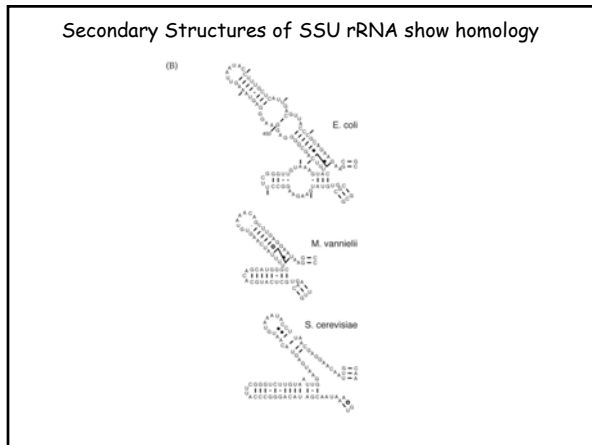
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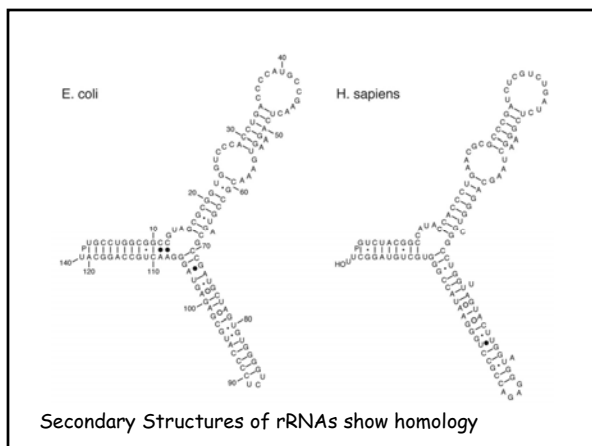
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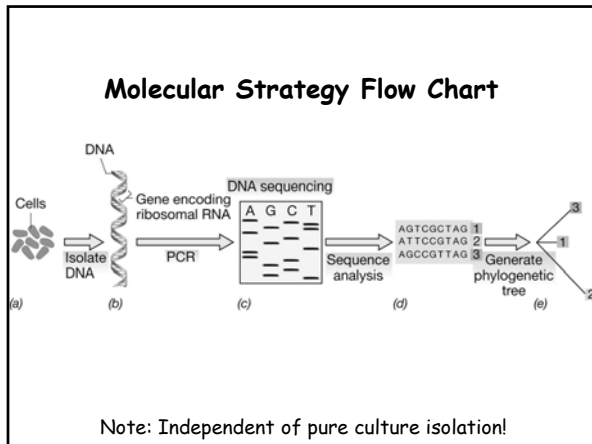
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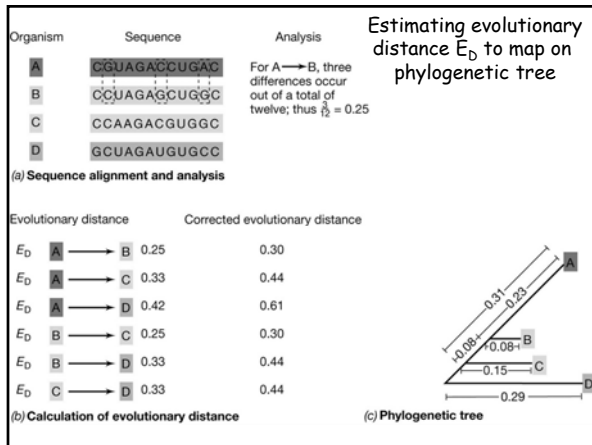
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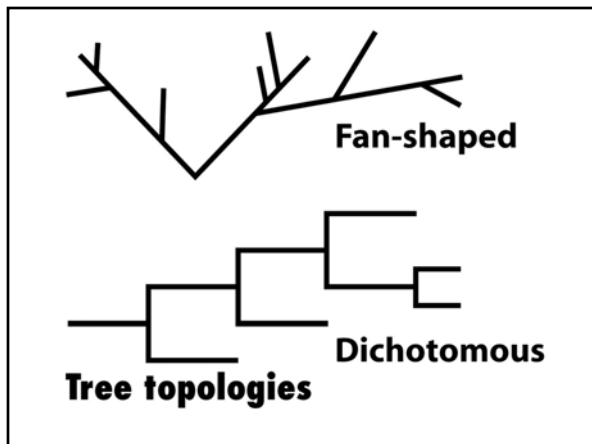
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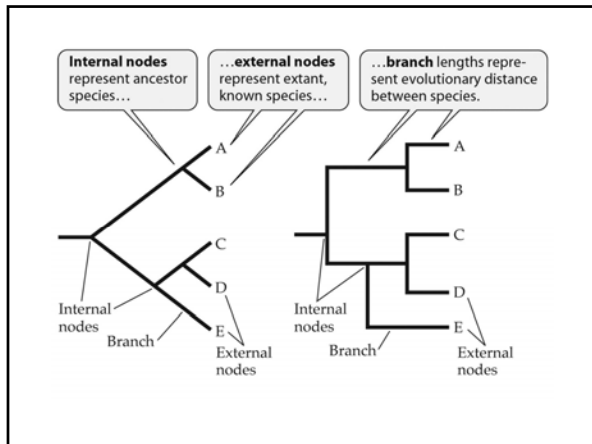
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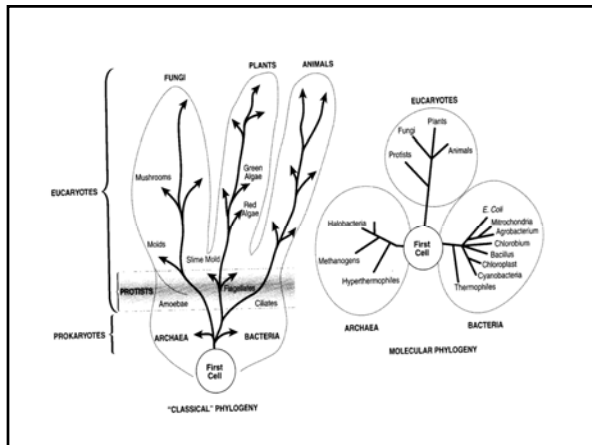
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**Table 11.1** Signature sequences from 16S or 18S rRNA defining the three domains of life

Oligonucleotide signatures <sup>a</sup>	Approximate position <sup>b</sup>	Occurrence among <sup>c</sup>		
		Archaea	Bacteria	Eukarya
CACYYG	315	0	>95	0
AAACUAAA	910	3	100	0
AAACUAAAG	910	100	0	100
YUYAAUUG	960	100	<1	100
CAACCYCR	1110	0	>95	0
UCCUG	1380	>95	0	100
UACACACCG	1400	0	>99	100
CACACACCC	1400	100	0	0

<sup>a</sup> Y, any pyrimidine; R, any purine.  
<sup>b</sup> Refer to Figure 11.11c for numbering scheme of 16S rRNA.  
<sup>c</sup> Occurrence refers to percentage of organisms examined in any domain that contain that sequence.

Signature sequences can be obtained at any level of taxonomic hierarchy

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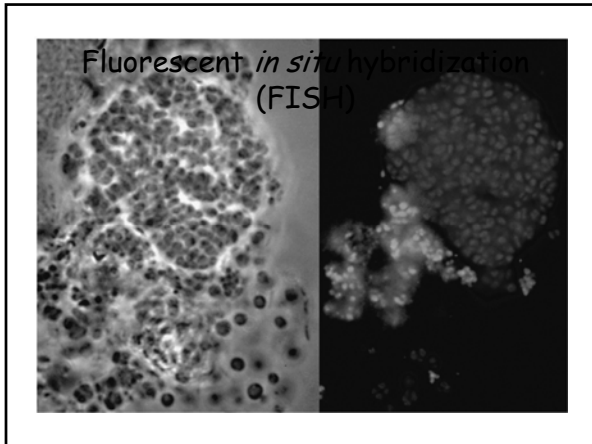
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### Take Home Message

- Phylogeny is right or wrong, we try to infer it the best we can.
- Taxonomy is useful or not, depending upon your point of view.
- Phylogeny allows us to ask testable questions, e.g., hypothesis testing.
  - microbial ecology relationships can now be truly examined
  - relationships between MOs and their genes can be studied
  - infer dynamics of sequence change (Rolex vs Timex)

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### Inferring evolutionary relationships from a phylogenetic tree

- Key word is inference (not always correct!)
- Some lineages accumulate mutations faster than others - generation times and selective pressures differ.
- So, molecular clocks are distorted ("soft watches").
- For this reason, mutation frequency cannot be calibrated to units of time. Tree can be calibrated to fossil record or geological evidence.

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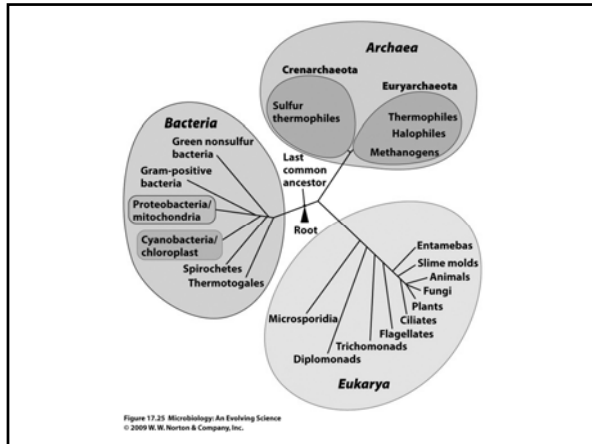
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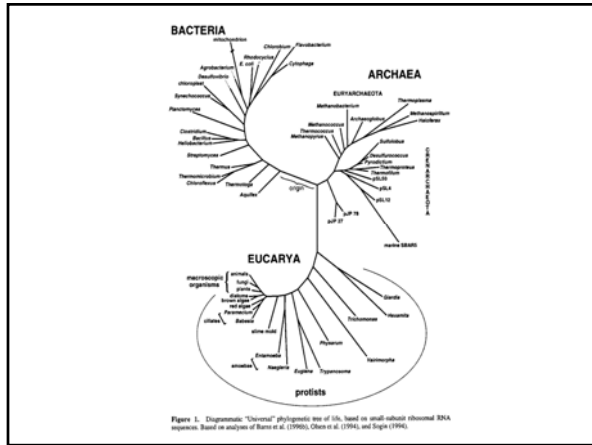
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**Some Lessons from the BIG TREE: Map of the Biological Record**

Single origin for all life on Earth...

- Central Dogma intact.
- ATP and PMF are universal themes.
- Uniformity among chiral carbon compounds (sugars & AAs).
- Hot start origin...

General topology implies:

- Three "primary lines of evolutionary descent."
- The Eucarya "nuclear" lineage almost as old as other two.
- Prokaryotes split between *Bacteria* and *Archaea*.
- Shown for only a limited number of representative org's.
- Mitochondria and chloroplasts proven to be of bacterial origin.

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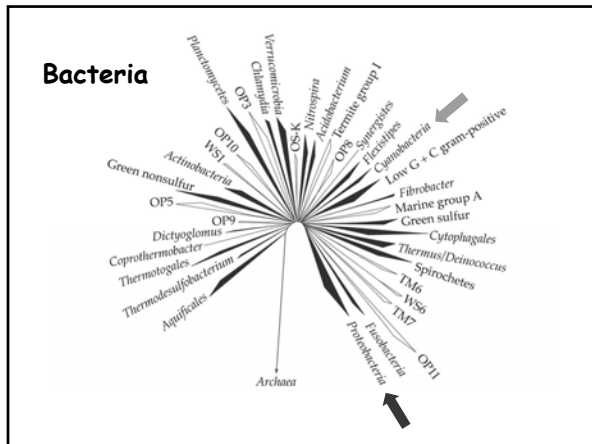
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**Some Lessons from the BIG TREE: Map of the Biological Record**

Evolutionary "clock" is NOT constant between different lineages

- Terminal nodes NOT all the same length, so not constant for all organisms either!
- Endosymbionts sped up very fast (semi-autonomous)
- Eucarya – Fast clocks
- Archaea – Slow clocks
- Bacteria – Intermediate

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**Table 17.3 Three domains of life.**

Characteristic	Traits of living organisms		
	All cells on Earth resemble each other		
Chromosomal material	Double-stranded DNA		
RNA transcription	Common ancestral RNA polymerase		
Translation	Common ancestral rRNAs and elongation factors		
Protein	Common ancestral functional domains		
Cell structure	Aqueous cell compartment bounded by a membrane		
	Comparison of domains		
	Bacteria	Archaea	Eukaryotes
	Archaea resemble bacteria		
Cell volume	1–100 $\mu\text{m}^3$ (usually)		1–10 <sup>6</sup> $\mu\text{m}^3$
DNA chromosome	Circular (usually)		Linear
DNA organization	Nucleoid		Nucleus with membrane
Gene organization	Multigene operons		Single genes
Metabolism	Denitrification, N <sub>2</sub> fixation, lithotrophy, respiration, and fermentation		Respiration and fermentation
Multicellularity	Simple		Simple or complex

Table 17.3 part 1 Microbiology: An Evolving Science  
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**Some Lessons from the BIG TREE: Map of the Biological Record**

What does genome sequencing and study of functional genomics add to our perspective?

- The central information processing machinery encompasses core genome.
- Metabolic functions, that's when relationships get murky.
- Endosymbiosis involves more than simply organelles, i.e., two-way transfer of genes with most going to the nucleus.
- Mitochondria have been at it much longer than chloroplasts.

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**A Bit on the Evolution of Evolutionary Thought**

A. Prior to the late 19th century, the concept of evolution was on the **evolutionary ladder**. Thus, we still deal in "higher and lower" eucaryotes (I try not to use these terms – they are dumb), "missing links," and "primitive" organisms.

B. In its milieu, *E. coli* is as highly evolved as are we. *E. coli* is **simple** (~5x10<sup>9</sup> bp genome), we are **complex** (~3x10<sup>9</sup> bps); complexity has nothing to do with *evolutionary advancement*.

C. Lineages evolve by diversification, not *progression*. !!!

D. There is no such thing as a *primitive* organism alive today. **Simple**, yes, but still a finely honed product of ~ 4 billion years under the selective hammer of the niches that it and its progenitors have occupied.

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**C-value paradox:  
Organism complexity  
does not correlate to  
genome size**

**TABLE 13.3 C values from eukaryotic organisms ranked by size**

Species	C value (kb)
<i>Nitzschia pediculus</i> (diatom)	35,000
<i>Drosophila melanogaster</i> (fruitfly)	180,000
<i>Paramecium aurelia</i> (ciliate)	190,000
<i>Gallus domesticus</i> (chicken)	1,200,000
<i>Ergasilphes cichoracearum</i> (fungus)	1,500,000
<i>Cyprinus carpio</i> (carp)	1,700,000
<i>Lampetra planeri</i> (lamprey)	1,900,000
<i>Bos constrictor</i> (snake)	2,100,000
<i>Parascaris equorum</i> (roundworm)	2,500,000
<i>Carcharias obscurus</i> (shark)	2,700,000
<i>Rattus norvegicus</i> (rat)	2,900,000
<i>Xenopus laevis</i> (toad)	3,100,000
<b>Homo sapiens (human)</b>	<b>3,400,000</b>
<i>Nicotiana tabacum</i> (tobacco)	3,800,000
<i>Paramecium caudatum</i> (ciliate)	8,600,000
<i>Schistocerca gregaria</i> (locust)	9,300,000
<i>Allium cepa</i> (onion)	18,000,000
<i>Coscinodiscus asteromphalus</i> (diatom)	25,000,000
<i>Lilium formosanum</i> (lily)	36,000,000
<i>Pinus resinosa</i> (pine)	68,000,000
<i>Amphiuma means</i> (newt)	84,000,000
<i>Protipterus aethiopicus</i> (lungfish)	140,000,000
<i>Ophioglossum petiolatum</i> (fern)	160,000,000
<i>Amoeba proteus</i> (amoeba)	290,000,000
<b>Amoeba dubia</b> (amoeba)	<b>670,000,000</b>

Compiled by Li and Graur (1991) from Cavalier-Smith (1985), Sparrow et al. (1972), and other references. The C value for humans is highlighted for reference.

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**Table 17.2** Comparison of *E. coli* and its primate host species<sup>a</sup>

Property	<i>E. coli</i>	<i>Homo sapiens</i>	Primates
Mol % G + C	48–52	42	42 <sup>b</sup>
16S–18S rRNA variability	>15 bases	?	<16 <sup>c</sup>
DNA/DNA reassociation	>70%	98.6% <sup>d</sup>	>70% <sup>e</sup>

<sup>a</sup>Adapted from J. T. Staley, *ASM News*, 1999.

<sup>b</sup>Value for all primates.

<sup>c</sup>Mouse 18S rRNA differs from humans by 16 bases.

<sup>d</sup>Comparison between *Homo sapiens* and chimpanzee.

<sup>e</sup>Comparison between *Homo sapiens* and lemurs.

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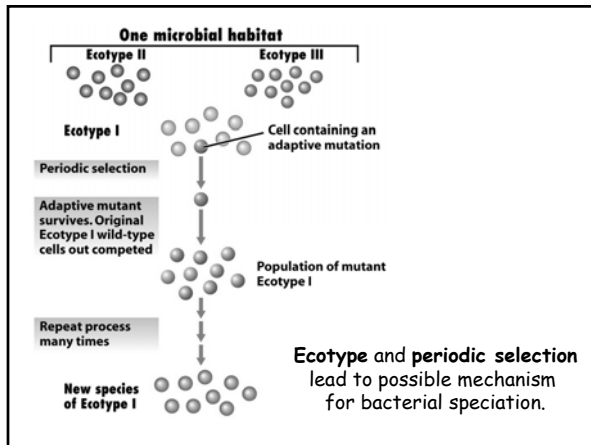
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