



Classifying microbes:

How do we sort all this out?

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.

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.

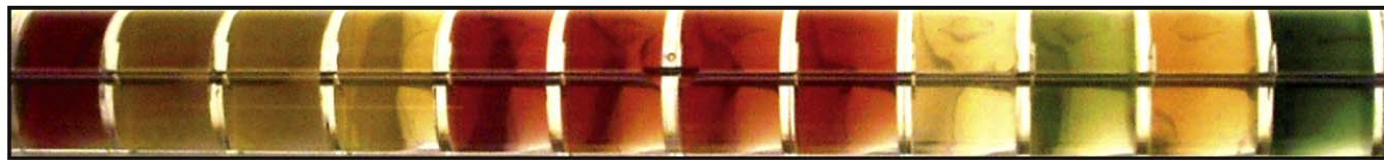
Slowly evolving molecules (e.g. rRNA) used for large-scale structure; "fast- clock" molecules for fine-structure.

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!)

Table 11.4 Some phenotypic characteristics of taxonomic value

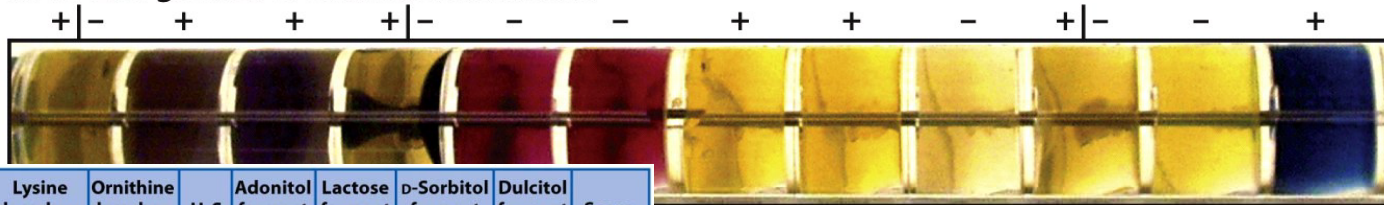
Major category	Components
I. Morphology	Shape; size; Gram reaction; arrangement of flagella, if present
II. Motility	Motile by flagella; motile by gliding; motile by gas vesicles; nonmotile
III. Nutrition and Physiology	Mechanism of energy conservation (phototroph, chemoorganotroph, chemolithotroph); 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	D-Glucose acid	Lysine decarbox.	Ornithine decarbox.	H ₂ S	Adonitol ferment	Lactose ferment	D-Sorbitol ferment	Dulcitol ferment	Score
Test Results:	+	+	+	+	-	-	+	-	
<i>Salmonella enterica</i>	0.99	0.98	0.97	0.95	0.01	0.01	0.95	0.96	3.3 E-2
<i>Proteus mirabilis</i>	0.99	0.01	0.99	0.98	0.01	0.02	0.01	0.01	9.2 E-5
<i>Yersinia enterocolitica</i>	0.99	0.01	0.95	0.01	0.01	0.05	0.99	0.01	8.7 E-5
<i>Enterobacter aerogenes</i>	0.99	0.98	0.98	0.01	0.98	0.95	0.99	0.05	8.9 E-6
<i>Klebsiella pneumoniae</i>	0.99	0.98	0.01	0.01	0.90	0.98	0.99	0.30	1.3 E-7

← Lactose
← Arabinose
← Sorbitol
← Voges Proskauer
← Dulcitol | phenylalanine
← Urea
← Citrate

Figure 17.19b Microbiology: An Evolving Science
Joan Slonczewski and BD Diagnostics

ID of an enteric bacterium

I. Isolation and microscopy

Isolation → Pure culture → Gram reaction/
morphology

II. General physiology

Gram-negative rod → Facultative → Ferments
lactose to
acid/gas

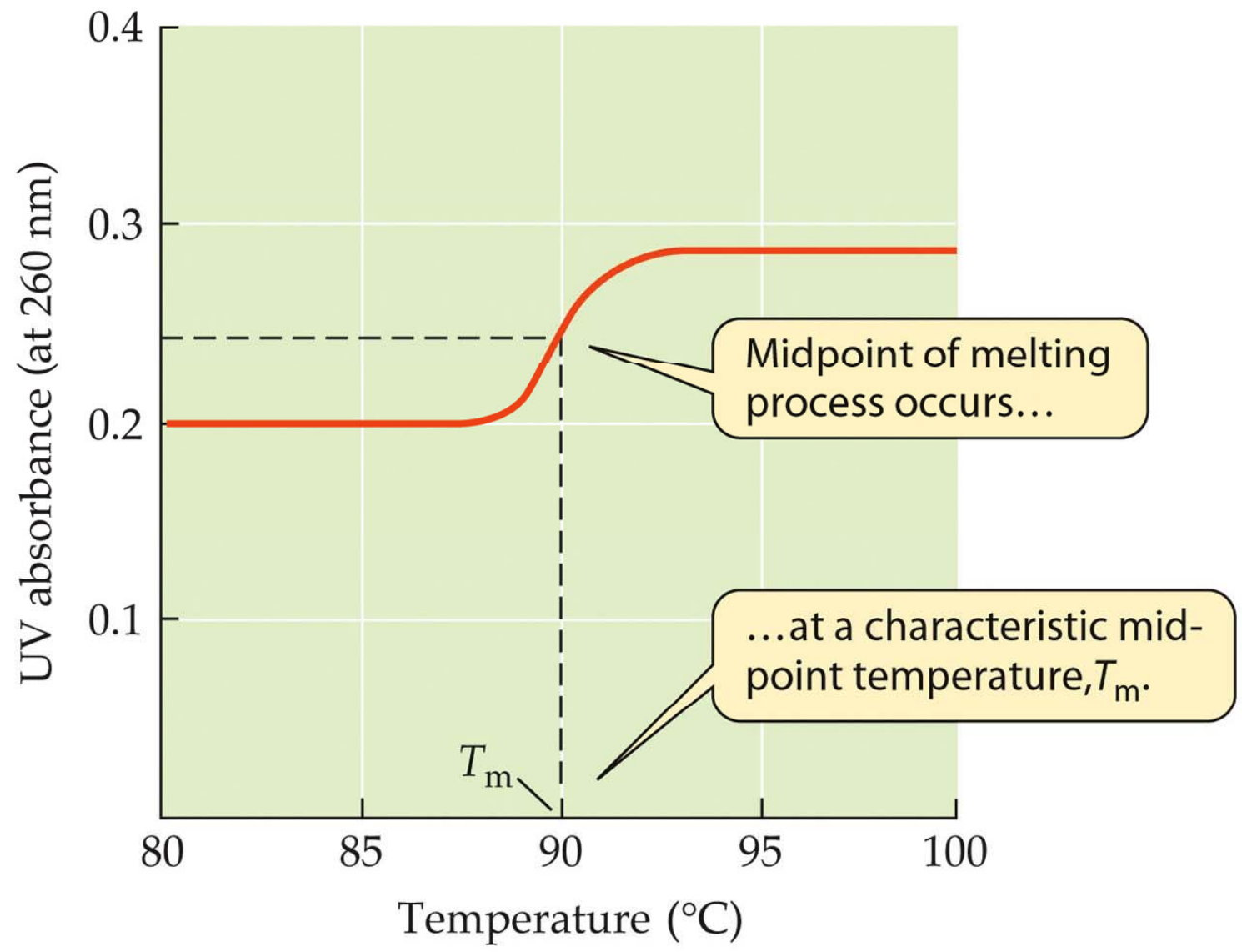
III. Detailed physiology

Facultative
lactose fermenter → Perform
series of
biochemical
tests → Positive:
indole, methyl red,
mucate;
Negative: citrate,
Voges-Proskauer,
H₂S

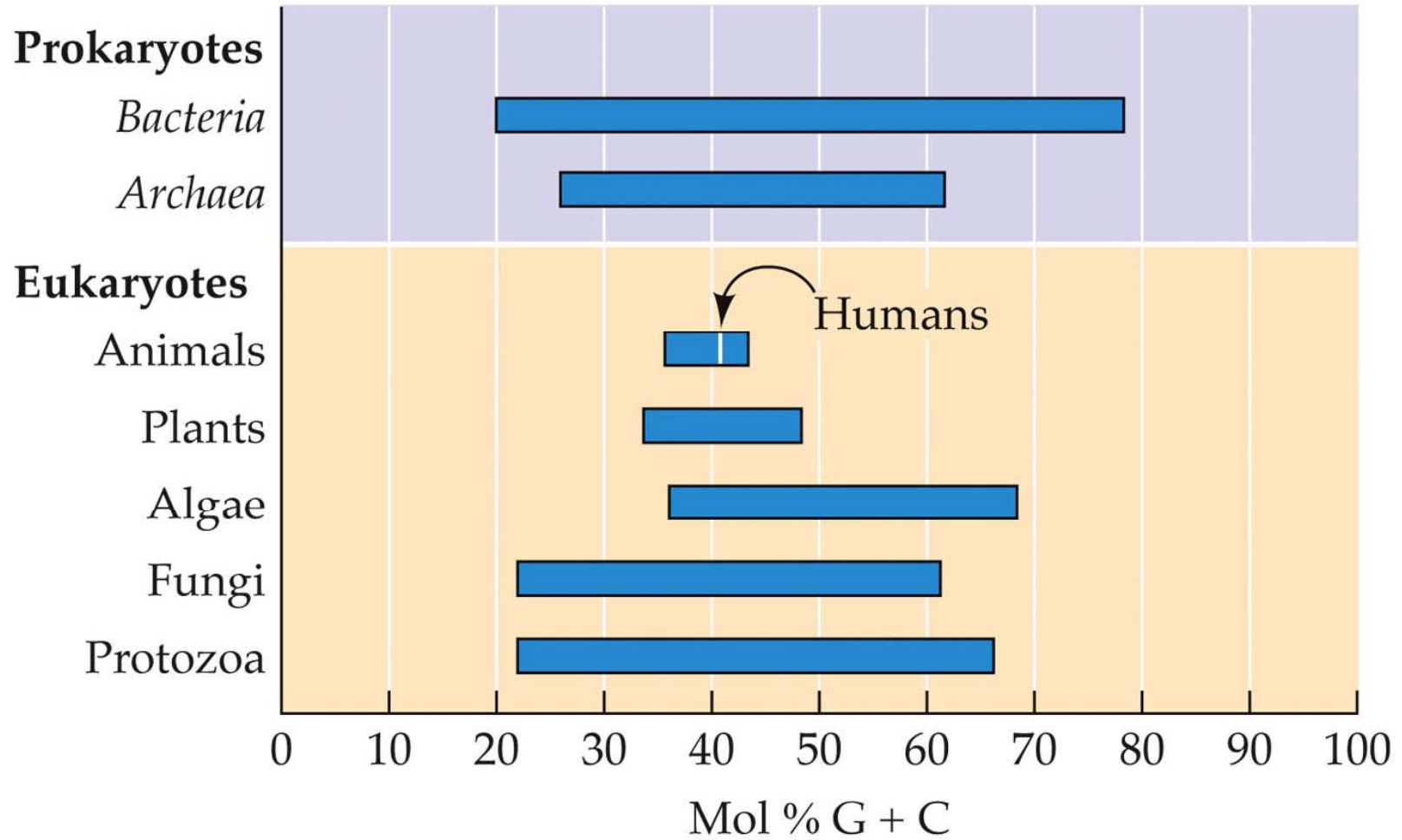
IV. Conclusion → *Escherichia coli*

Note: requires isolation in pure culture!

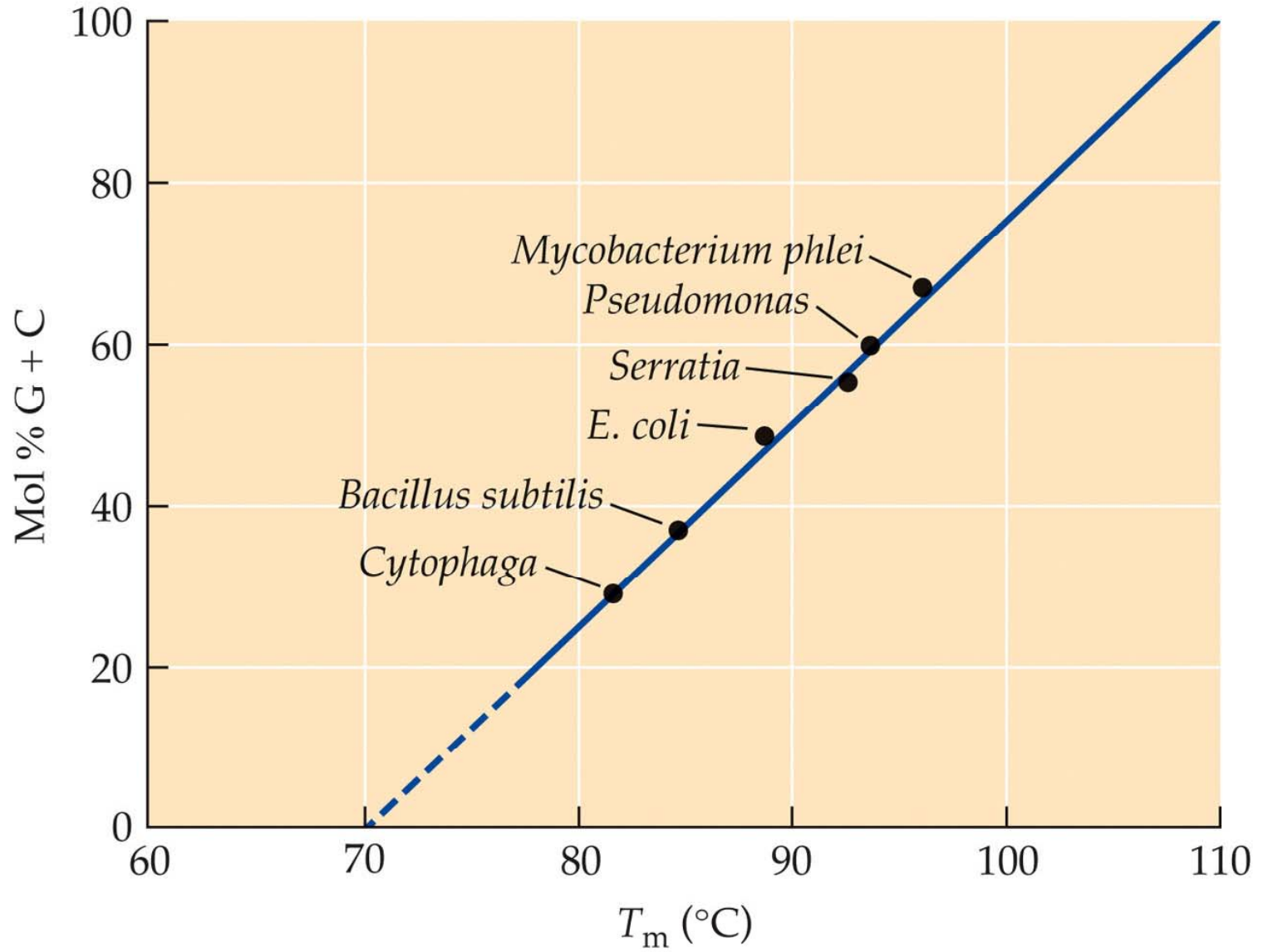
Hyperchromic Effect of DNA



Ranges of DNA base composition



G+C Ratios



DNA:DNA hybridization

Organisms to be compared:

Organism 1

Organism 2

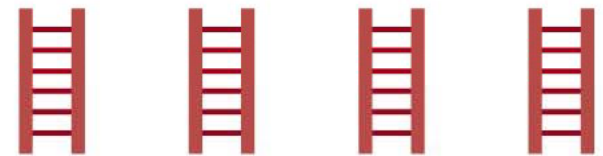
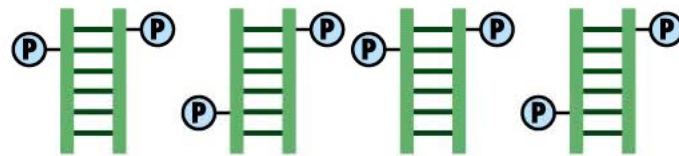
DNA preparation

DNA

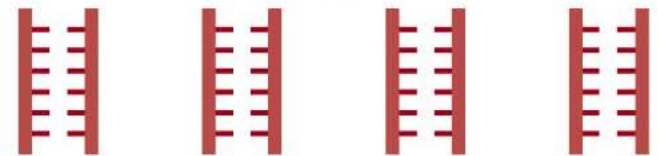
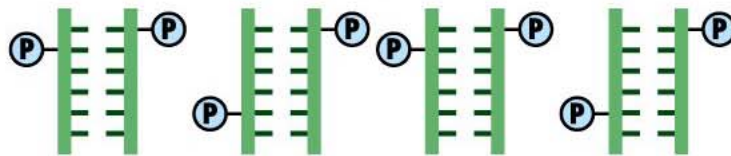
DNA

Shear and label (-P)

Shear DNA



Heat to denature

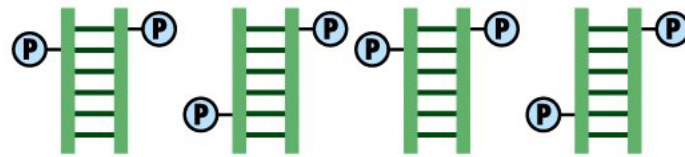


DNA:DNA hybridization

Hybridization experiment:

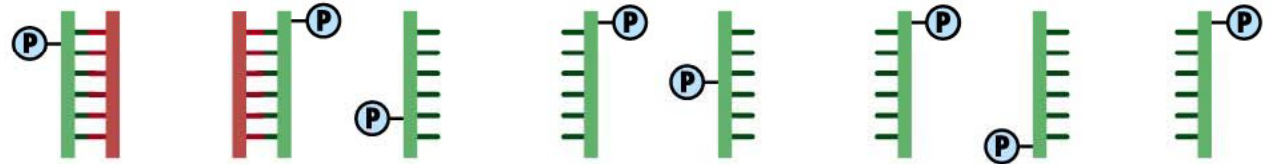
Mix DNA from two organisms—unlabeled DNA is added in excess:

1 x 1



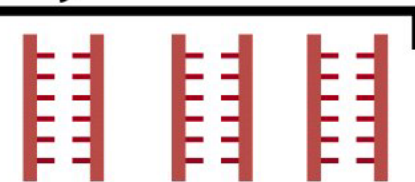
Hybridized DNA

1 x 2



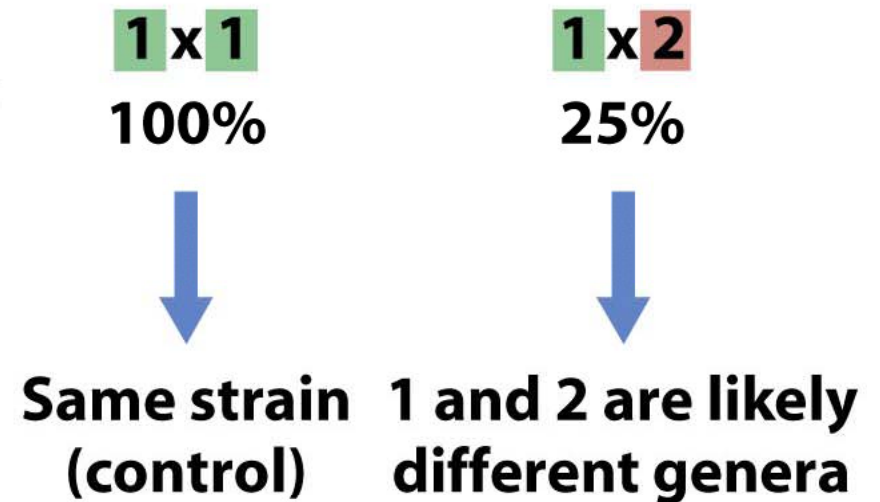
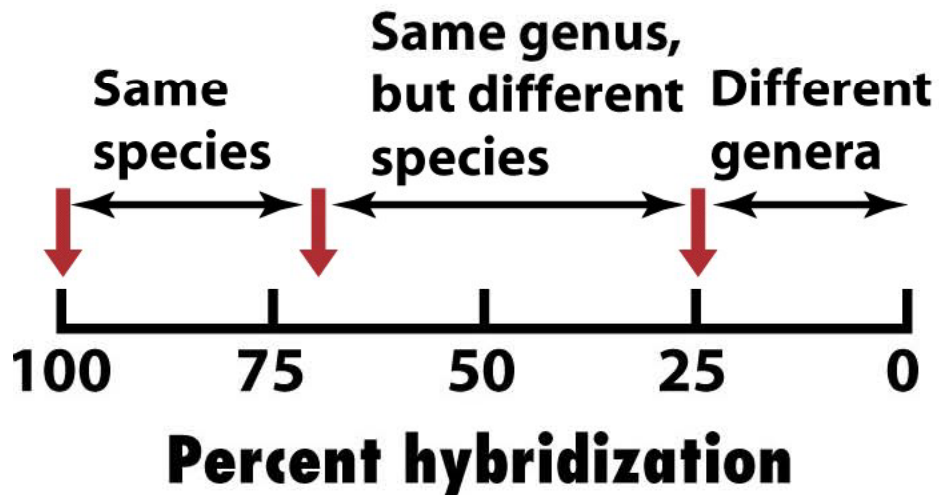
Hybridized DNA

Unhybridized DNA



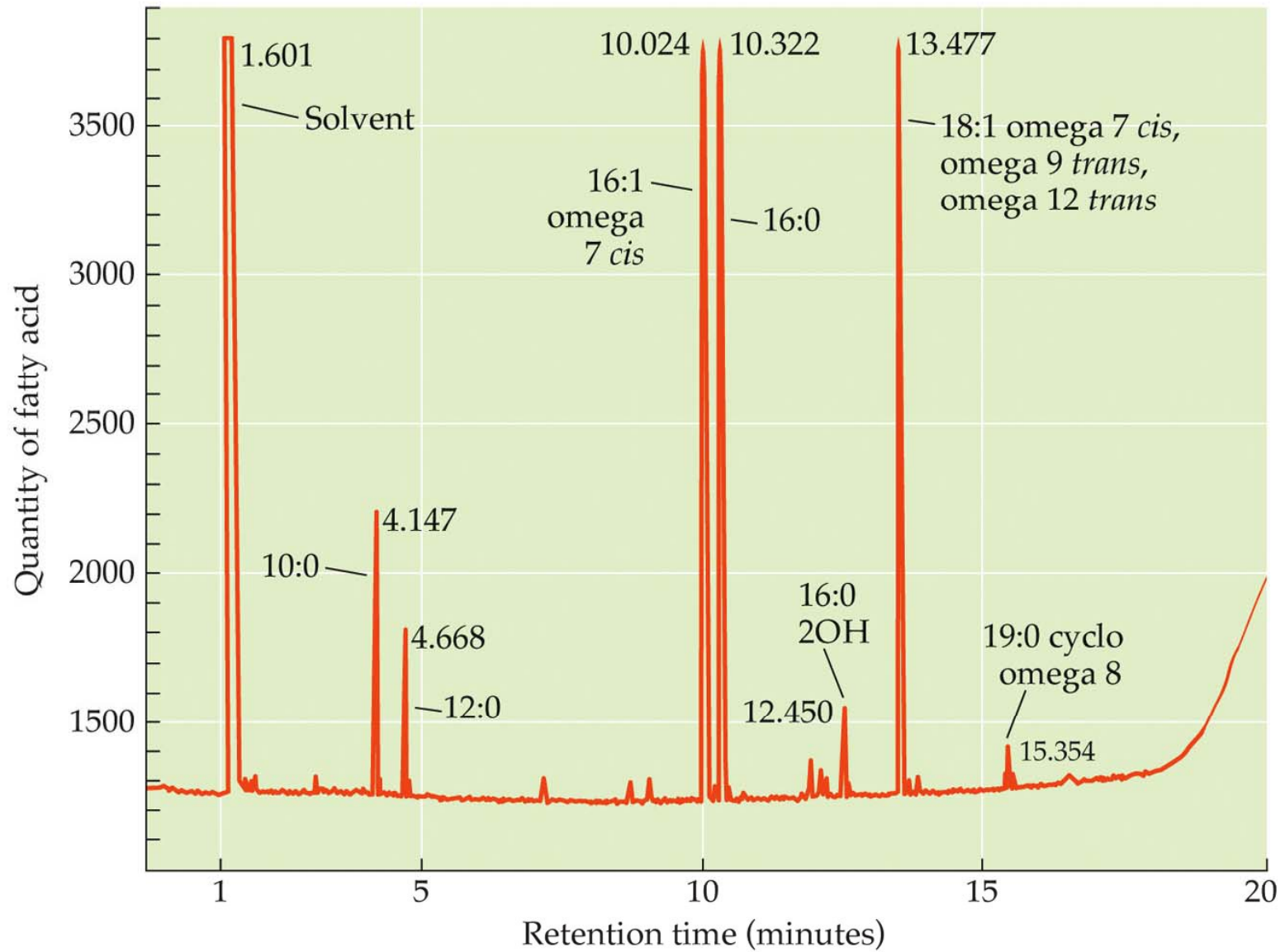
DNA:DNA hybridization

Results and interpretation:



70% or greater; considered same species

FAME analysis

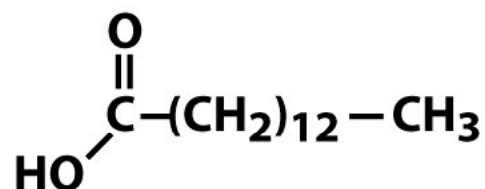


Classes of Fatty Acids in *Bacteria*

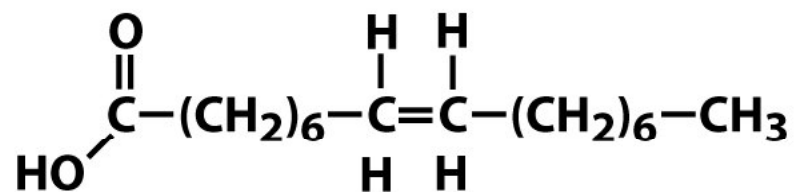
Class / Example

Structure of example

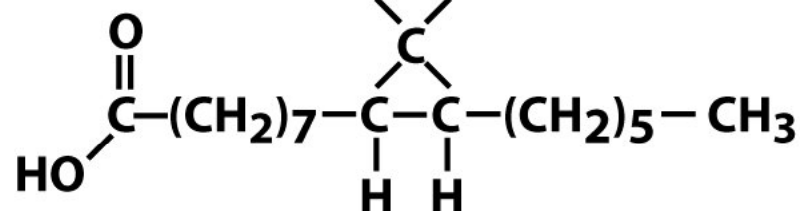
I. **Saturated:**
tetradecanoic acid



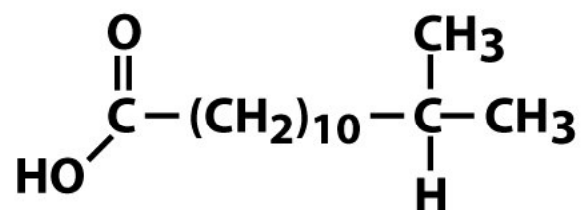
II. **Unsaturated:**
omega-7-cis
hexadecanoic acid



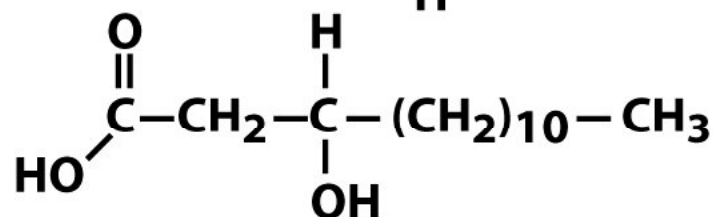
III. **Cyclopropane:**
cis 7, 8 methylene
hexadecanoic acid



IV. **Branched:**
13-methyltetradecanoic acid



V. **Hydroxy:**
3-hydroxytetradecanoic acid





Bacterial culture



Extract fatty acids



**Derivatize to form
methyl esters**



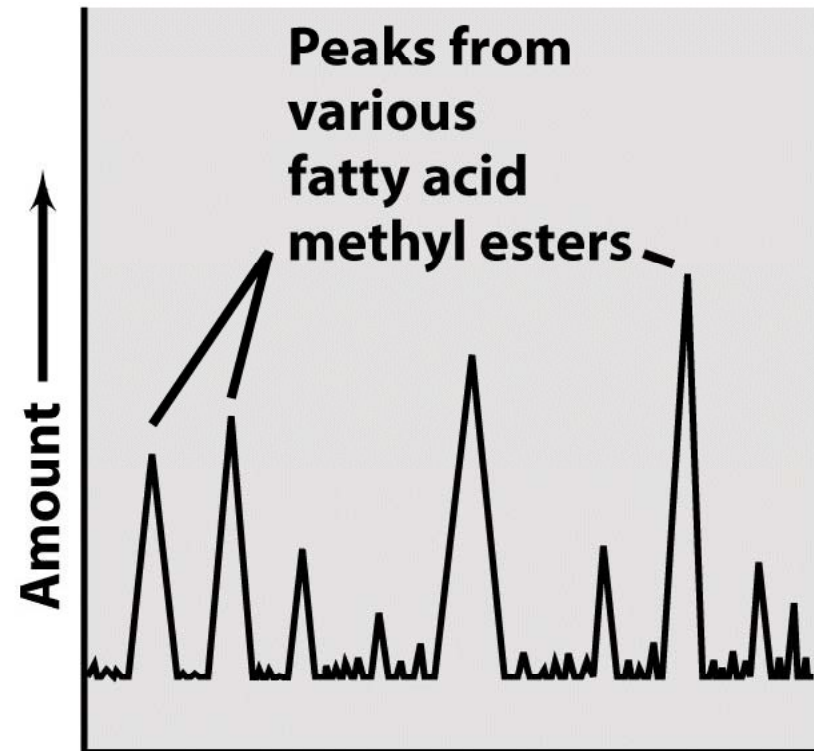
Gas chromatography



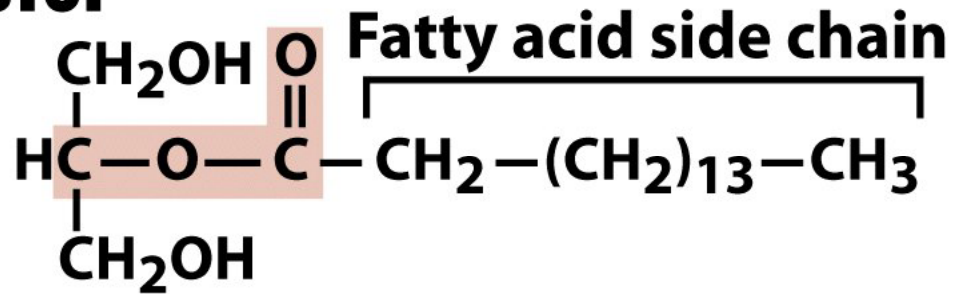
IDENTIFY ORGANISM



**Compare pattern of peaks
with patterns in database**

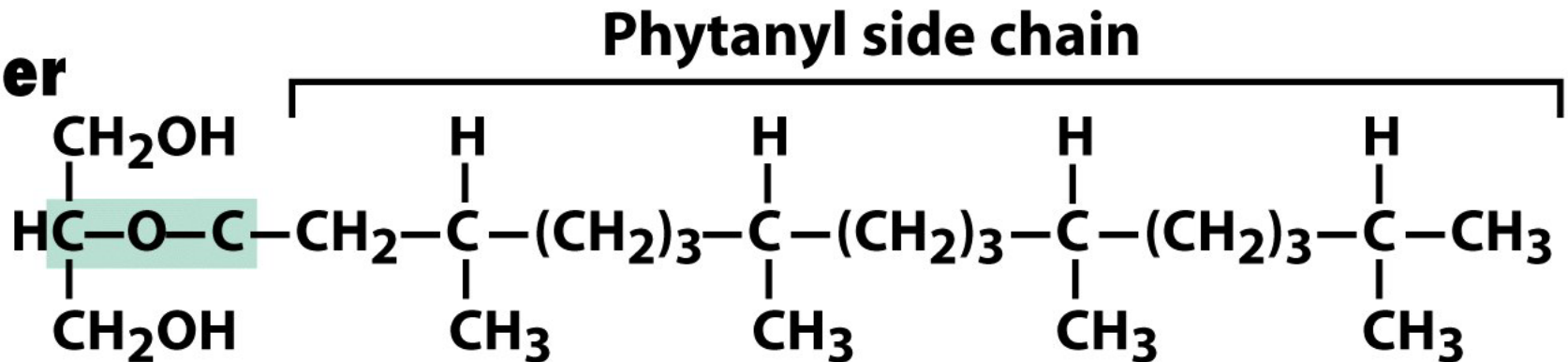


Ester



Bacteria, Eukarya

Ether



Archaea

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>

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	Streptomycineae		
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. jannaschii</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^a**

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

^a 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.



Bergey's Manual of
**Systematic
Bacteriology**



SECOND EDITION

Volume One

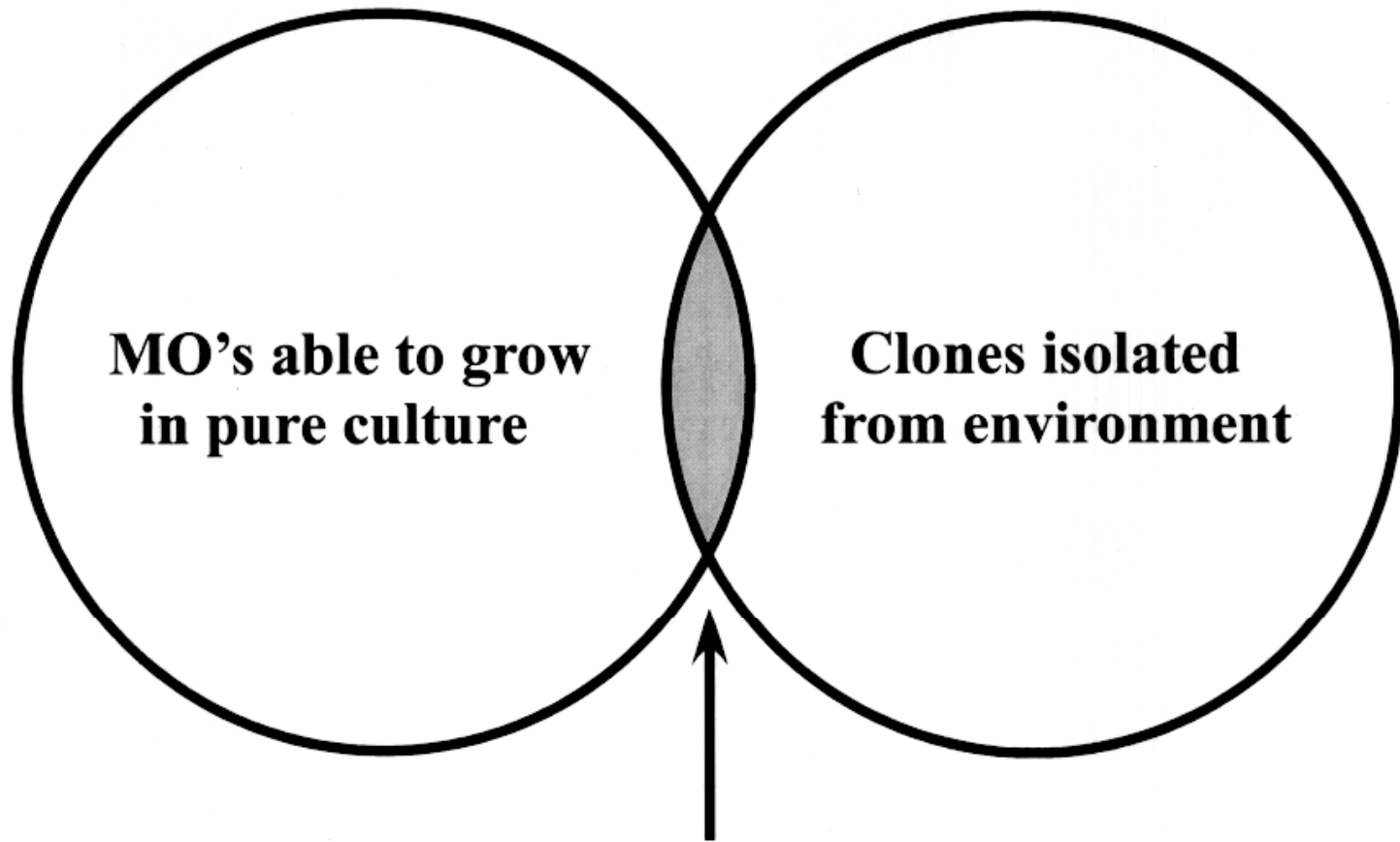
The *Archaea* and the Deeply Branching
and Phototrophic *Bacteria*

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.



**>1% Crossover
between these groups**

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.



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.

Regarding Molecular Phylogeny

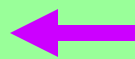
The Root of the Problem:

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

Molecular Phylogeny



Figure 17.25 (inset) Microbiology: An Evolving Science
Courtesy of Carl Woese



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


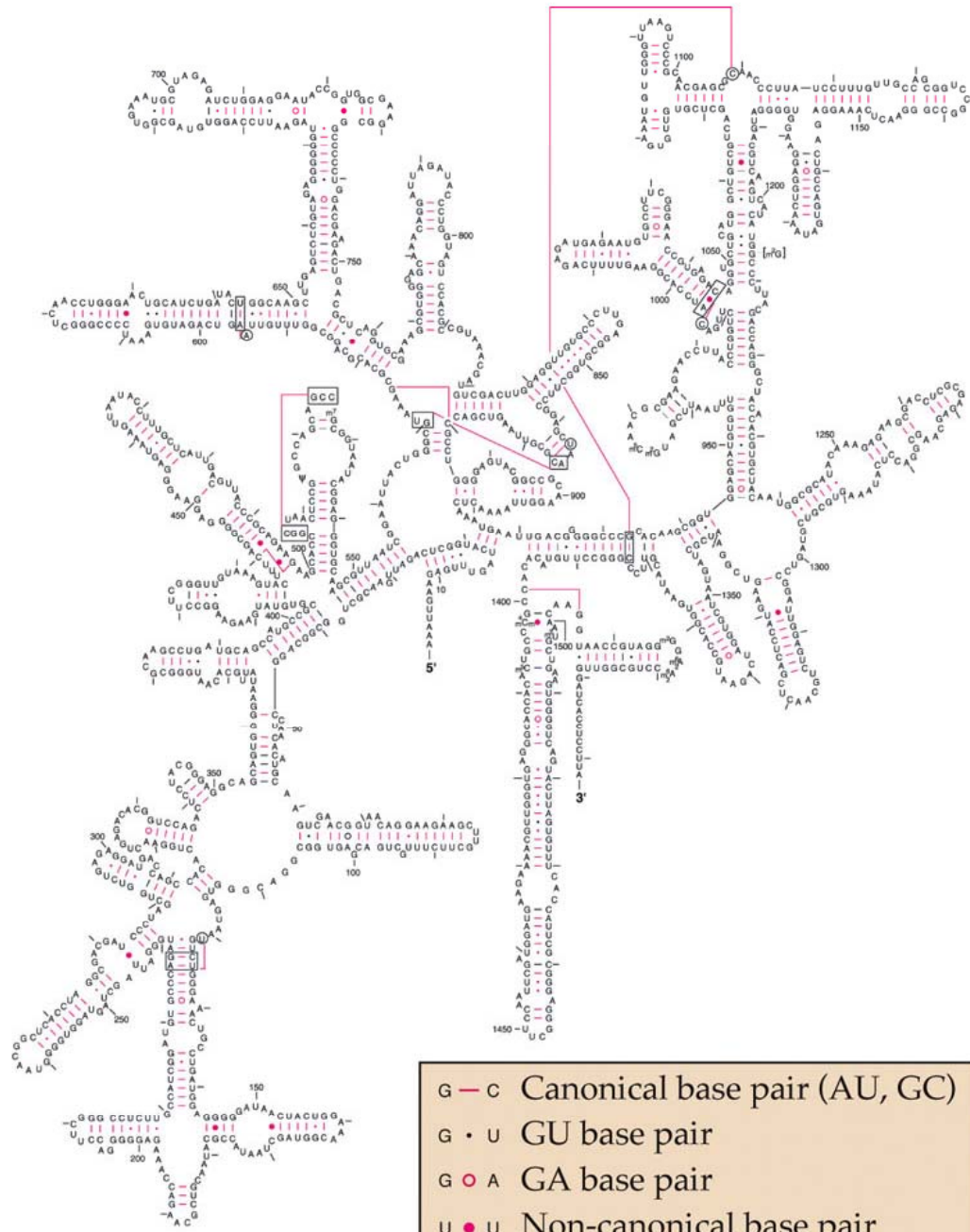
Pace (1984): Applied rRNA  concept to microbial ecology (census without culturing).



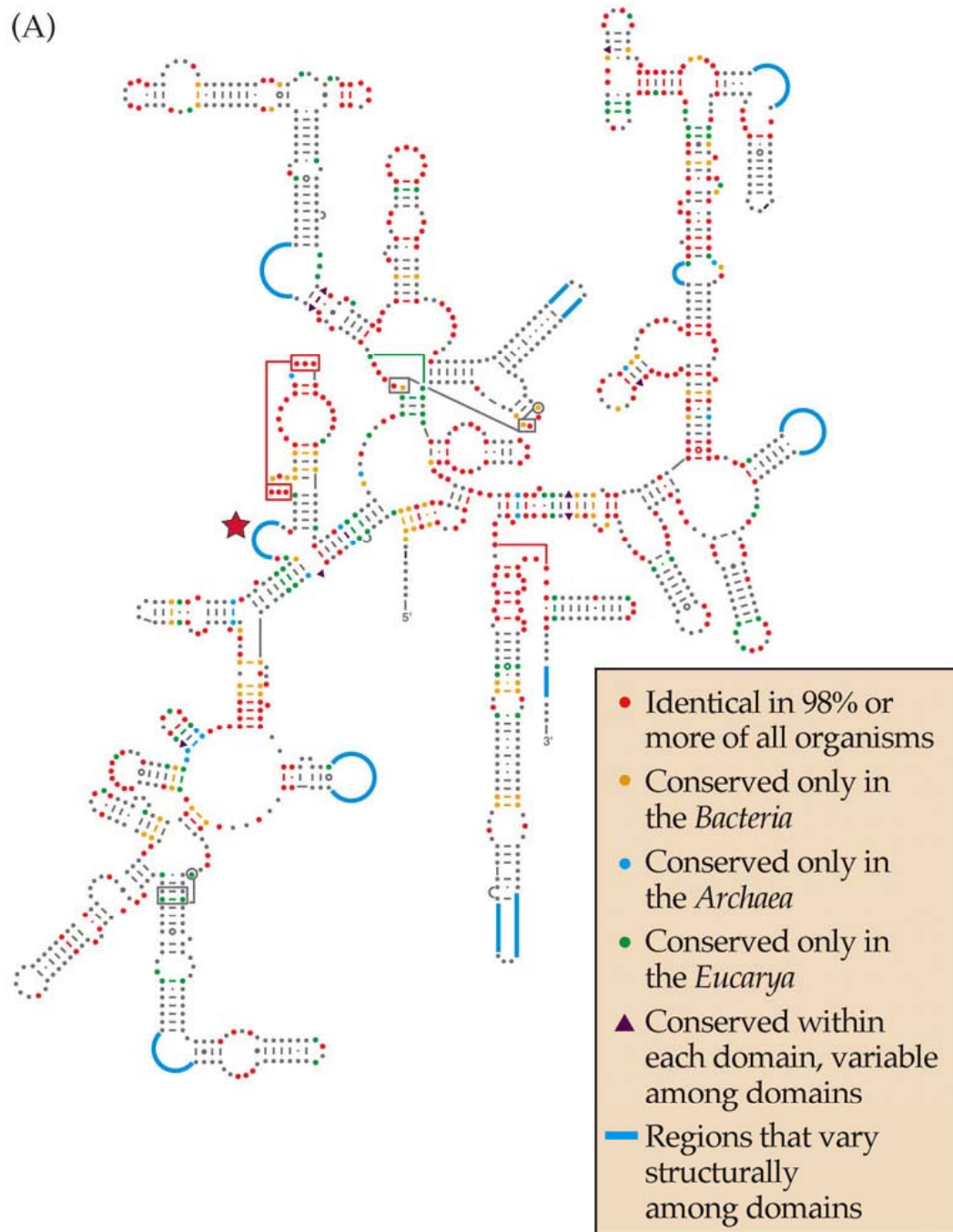
Figure 17.23a Microbiology: An Evolving Science
Courtesy of Norman R. Pace

Why ribosomal RNAs?

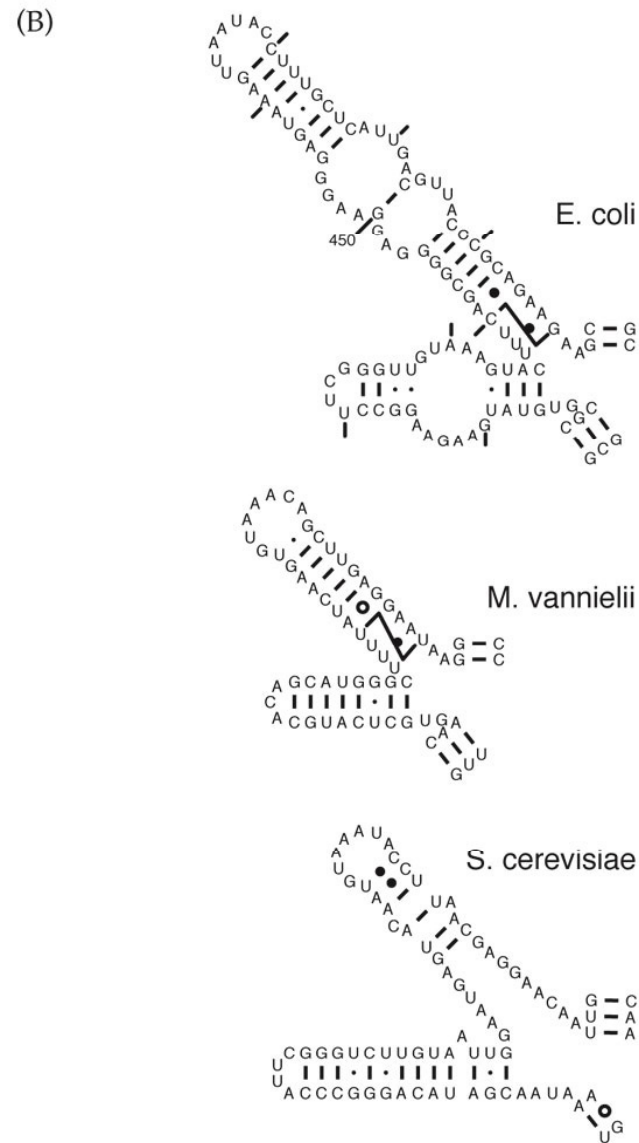
- Found among all living organisms (for 3.8 of the last 4.5 billion years). Integral part of protein synthesis machinery.
- Cell component analyses provide culture-independent means of investigating questions in microbial ecology (lack of morphology).
- rRNAs offer a type of sequence information that 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 $\sim 2.6 \times 10^6$ SSU rRNAs.



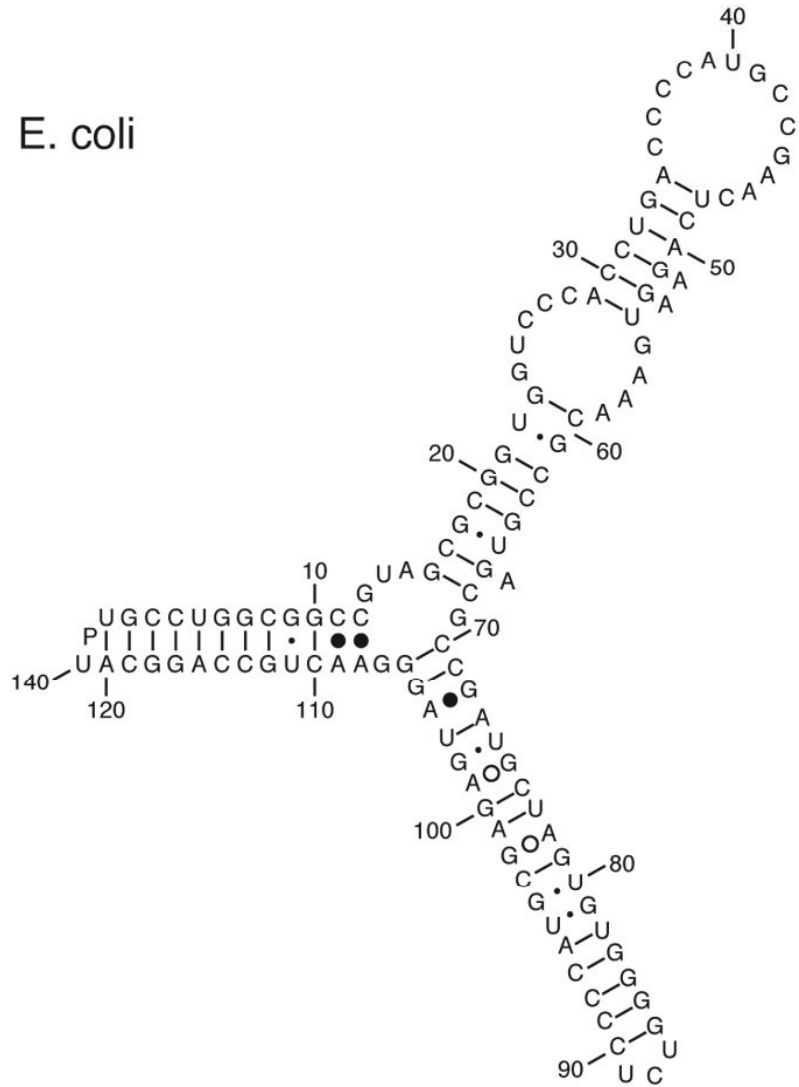
(A)



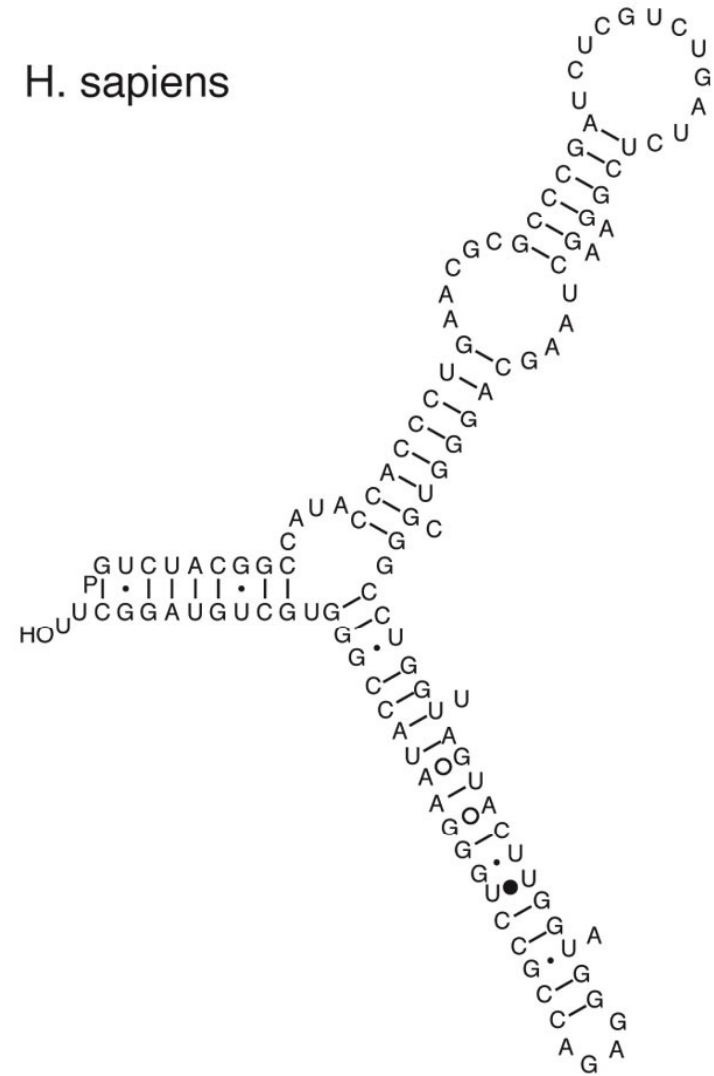
Secondary Structures of SSU rRNA show homology



E. coli

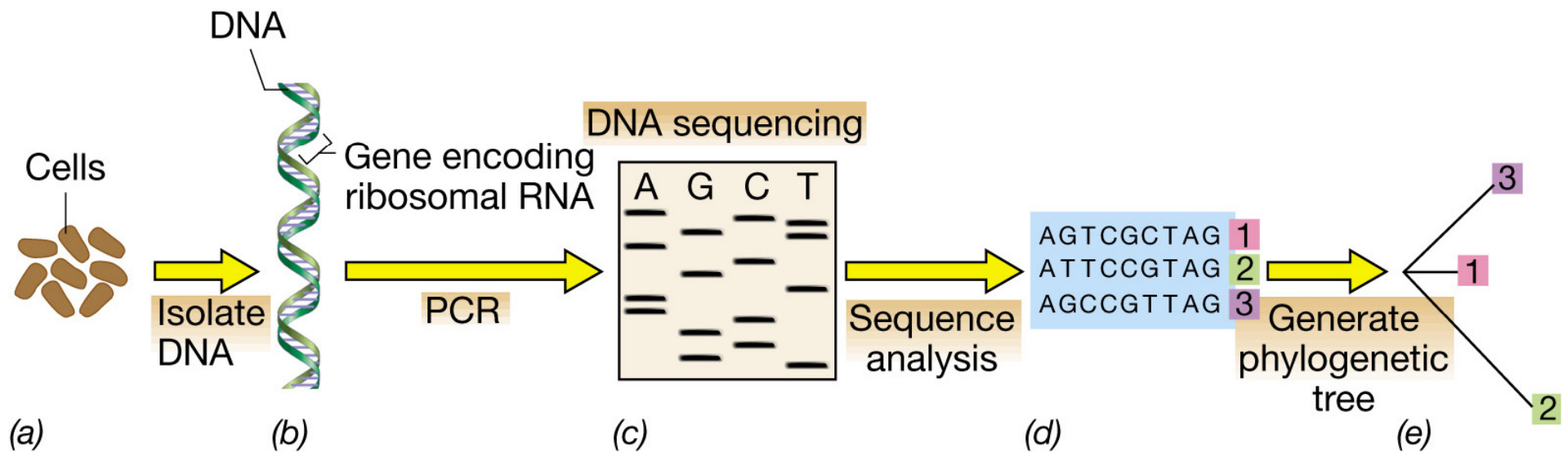


H. sapiens



Secondary Structures of rRNAs show homology

Molecular Strategy Flow Chart



Note: Independent of pure culture isolation!

Organism	Sequence
A	C G U A G A C C U G A C
B	C C U A G A G C U G G C
C	C C A A G A C G U G G C
D	G C U A G A U G U G C C

Analysis

For A → B, three differences occur out of a total of twelve; thus $\frac{3}{12} = 0.25$

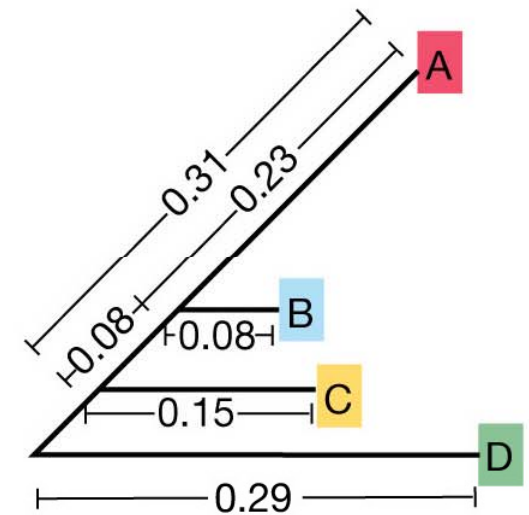
Estimating evolutionary distance E_D to map on phylogenetic tree

(a) Sequence alignment and analysis

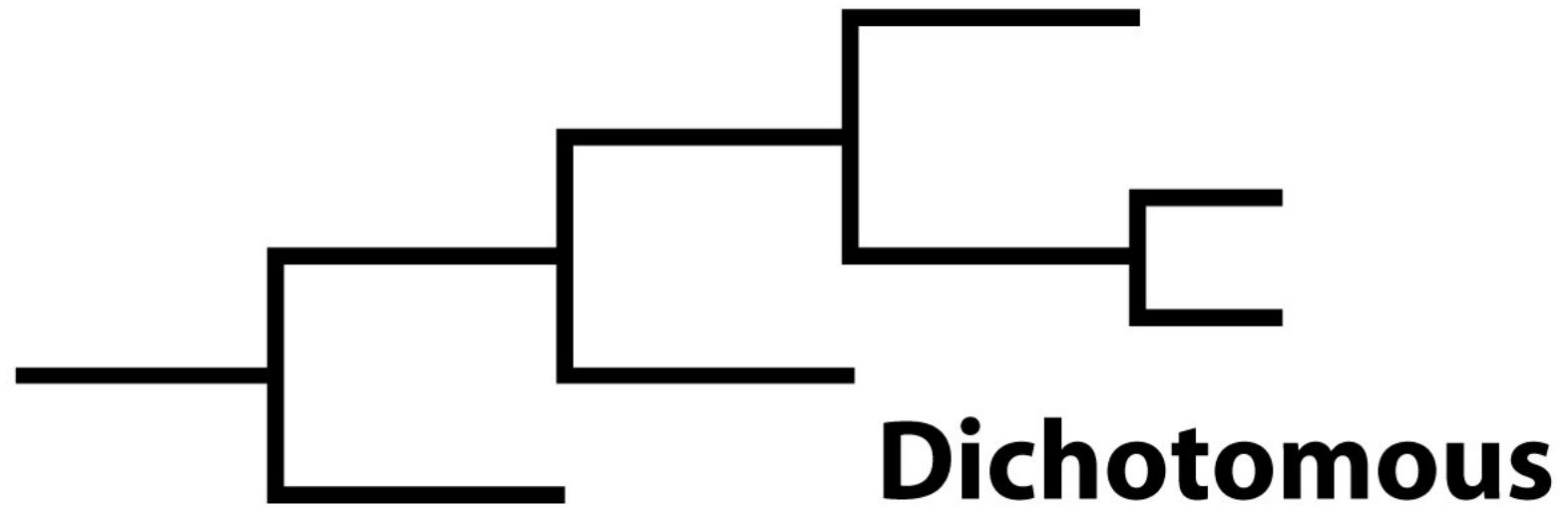
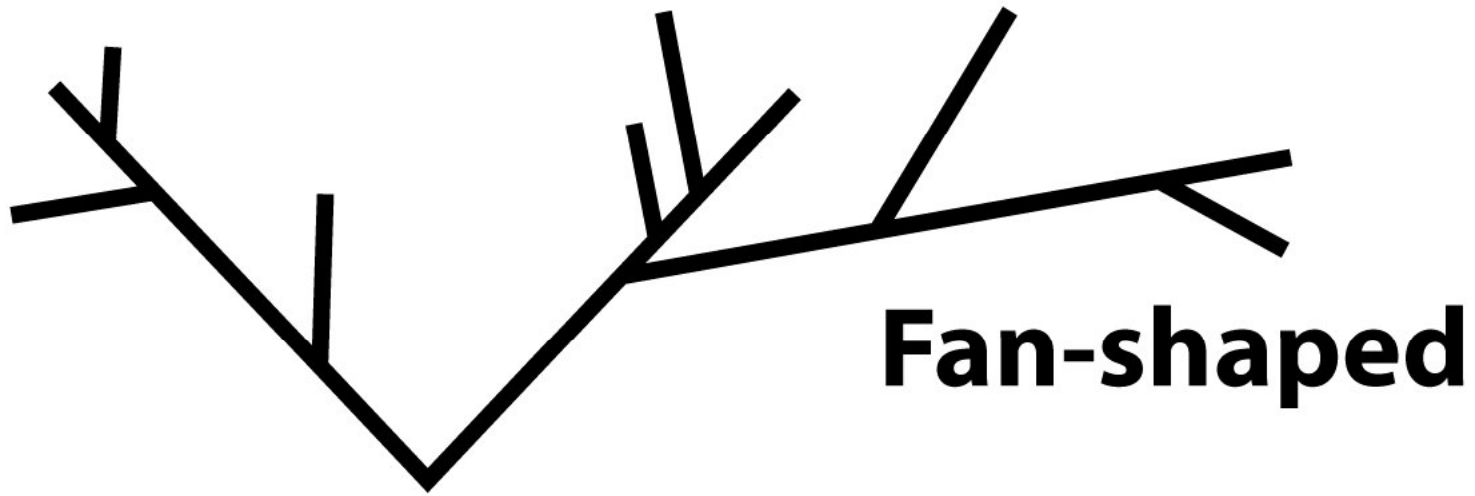
Evolutionary distance Corrected evolutionary distance

E_D	A → B	0.25	0.30
E_D	A → C	0.33	0.44
E_D	A → D	0.42	0.61
E_D	B → C	0.25	0.30
E_D	B → D	0.33	0.44
E_D	C → D	0.33	0.44

(b) Calculation of evolutionary distance



(c) Phylogenetic tree

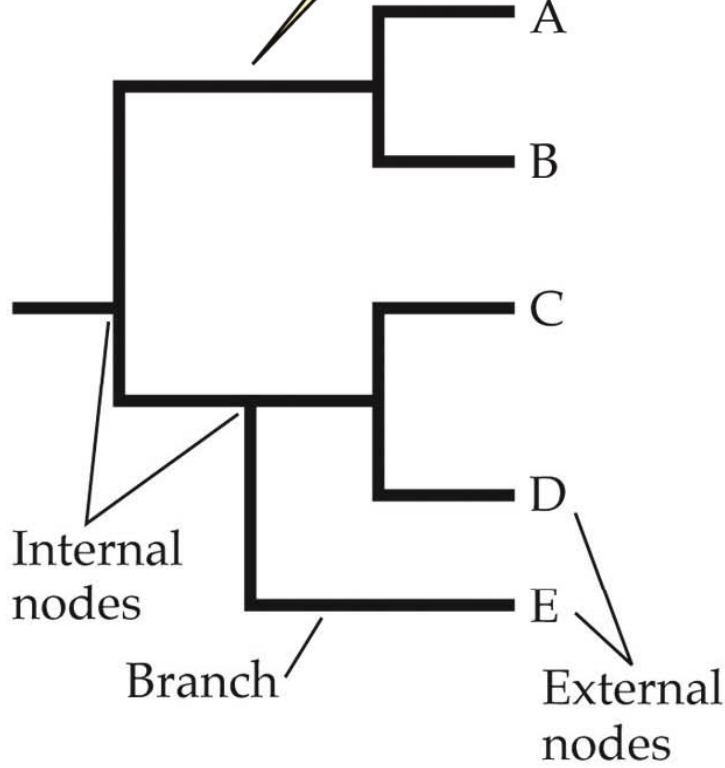
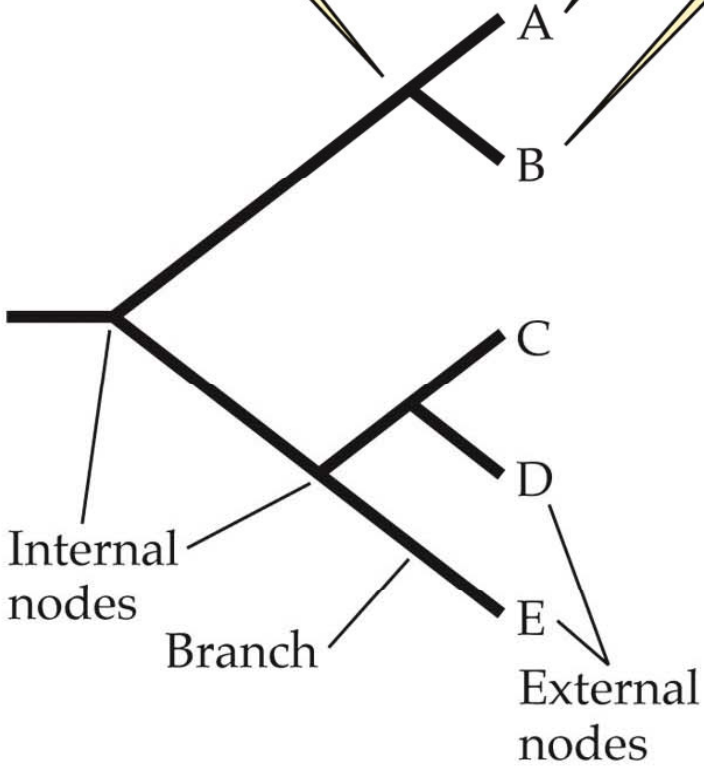


Tree topologies

Internal nodes
represent ancestor
species...

...**external nodes**
represent extant,
known species...

...**branch** lengths repre-
sent evolutionary distance
between species.



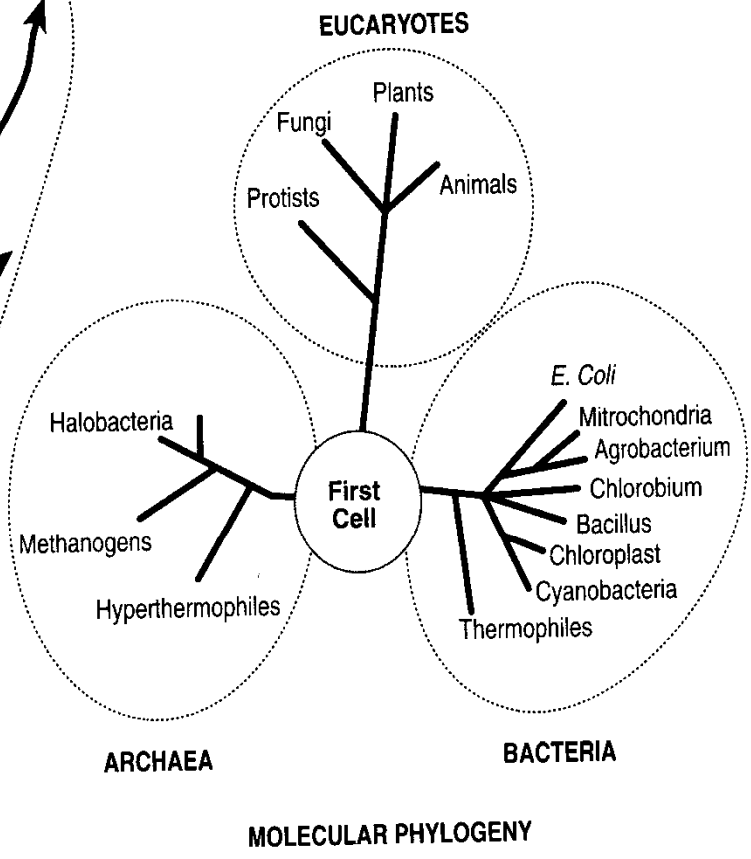
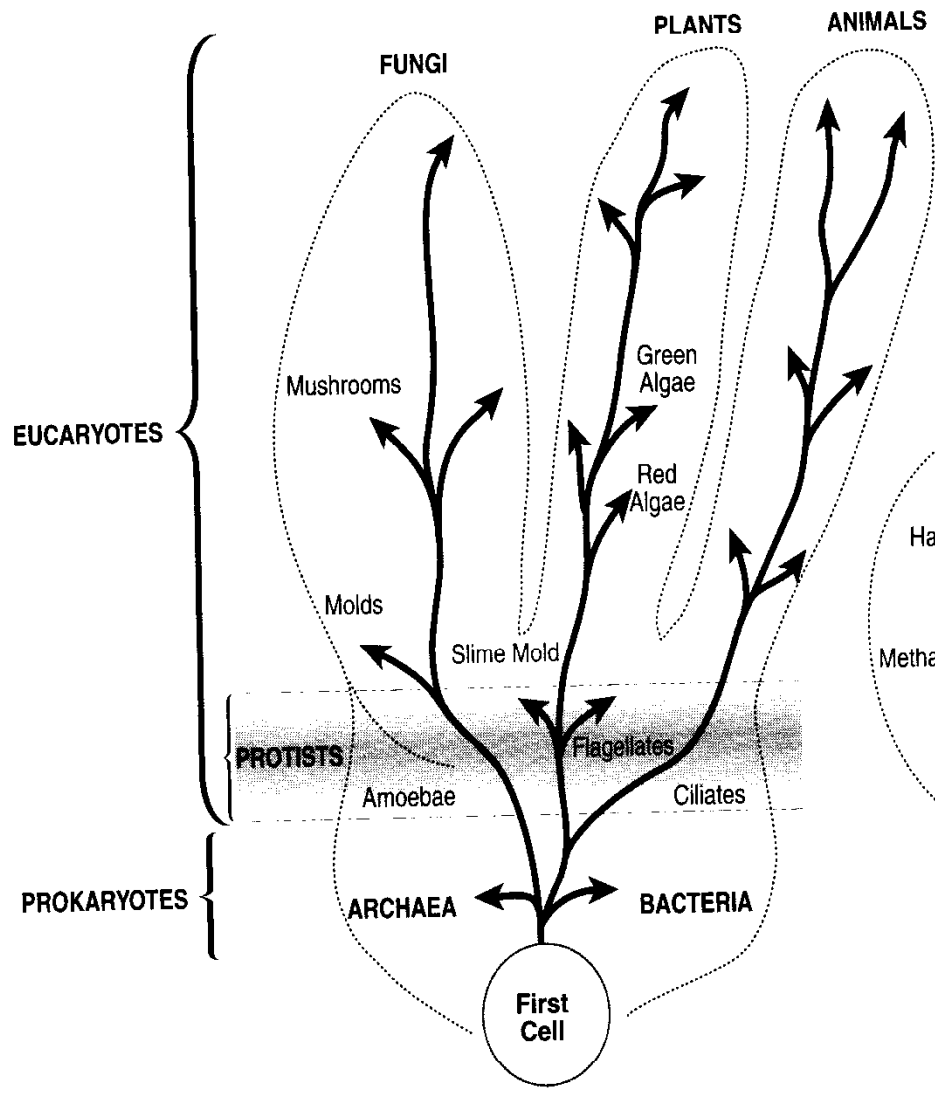


Table 11.1 Signature sequences from 16S or 18S rRNA defining the three domains of life

Oligonucleotide signatures^a	Approximate position^b	Occurrence among^c		
		Archaea	Bacteria	Eukarya
CACYYG	315	0	>95	0
AAACUCAAA	910	3	100	0
AAACUAAAAG	910	100	0	100
YUYAAUUG	960	100	<1	100
CAACCYYCR	1110	0	>95	0
UCCCUG	1380	>95	0	100
UACACACCG	1400	0	>99	100
CACACACCG	1400	100	0	0

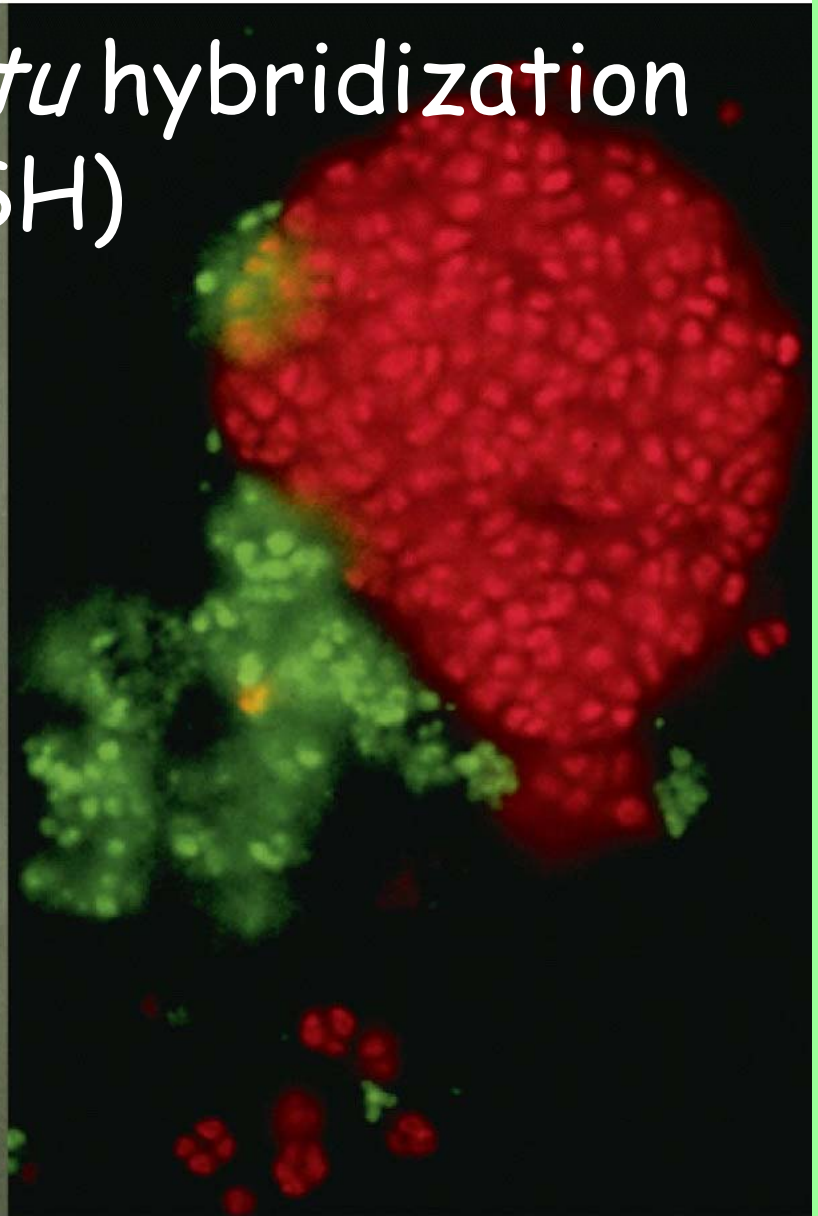
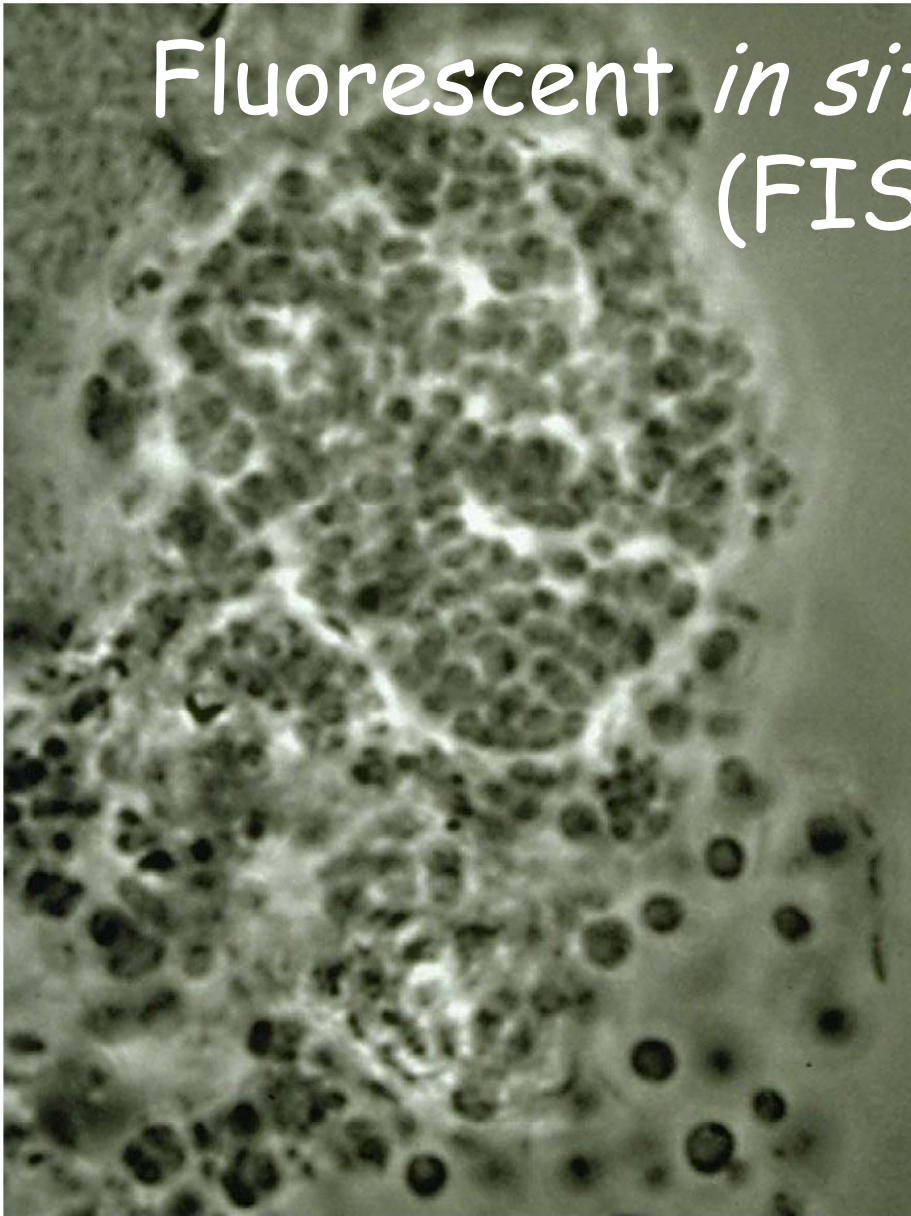
^a Y, any pyrimidine; R, any purine.

^b Refer to Figure 11.11c for numbering scheme of 16S rRNA.

^c 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

Fluorescent *in situ* hybridization
(FISH)



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)

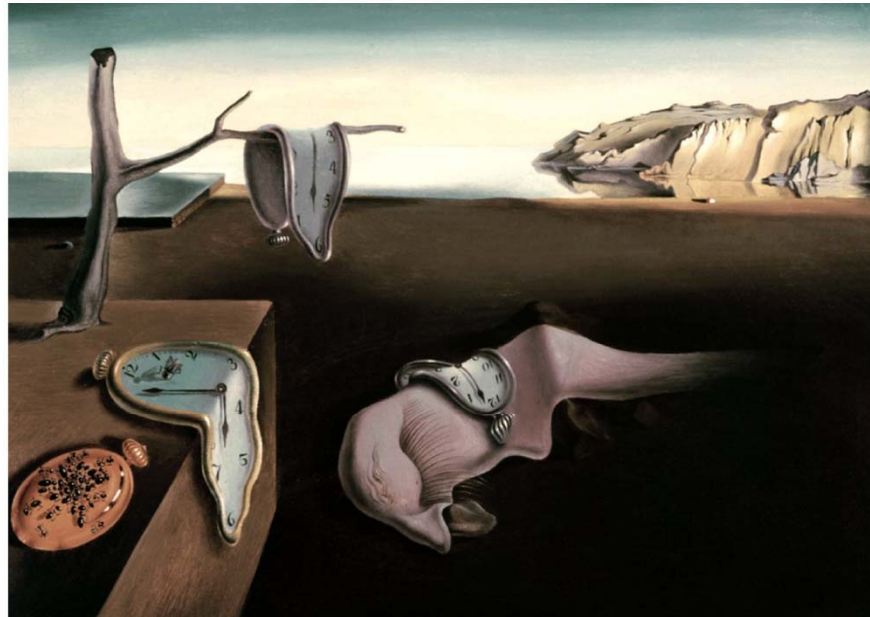
Inferring evolutionary relationships from a phylogenetic tree

-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.



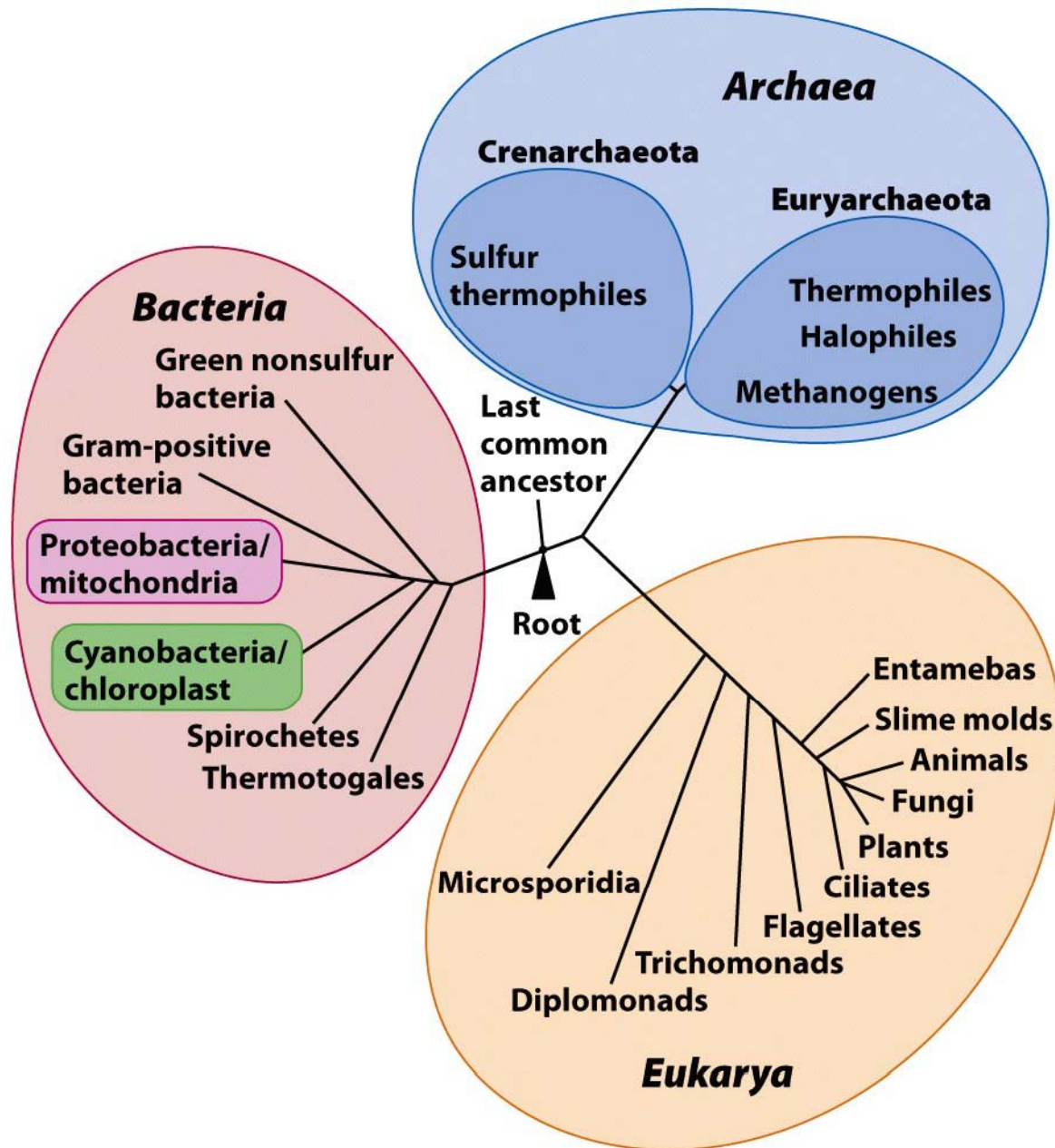


Figure 17.25 Microbiology: An Evolving Science
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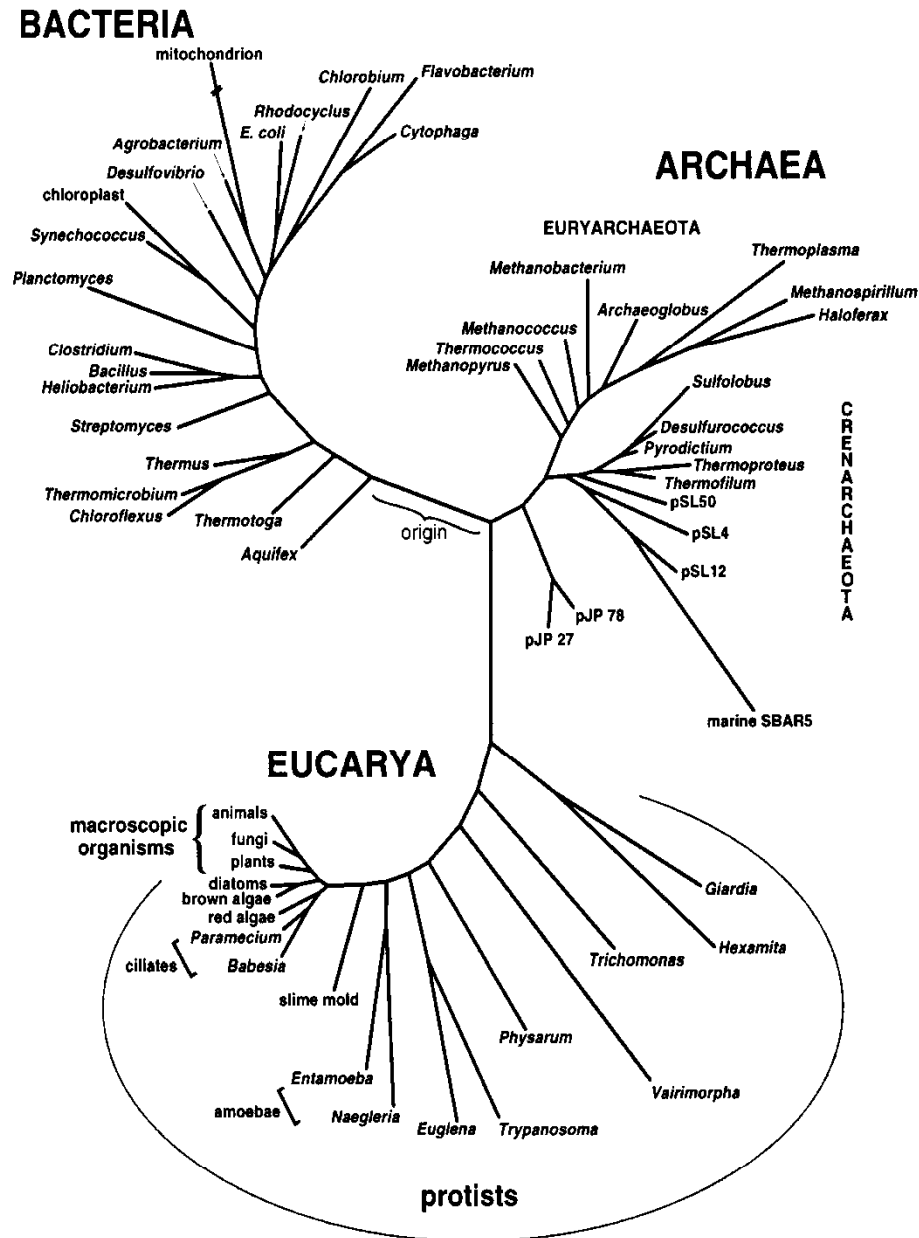


Figure 1. Diagrammatic "Universal" phylogenetic tree of life, based on small-subunit ribosomal RNA sequences. Based on analyses of Barns et al. (1996b), Olsen et al. (1994), and Sogin (1994).

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

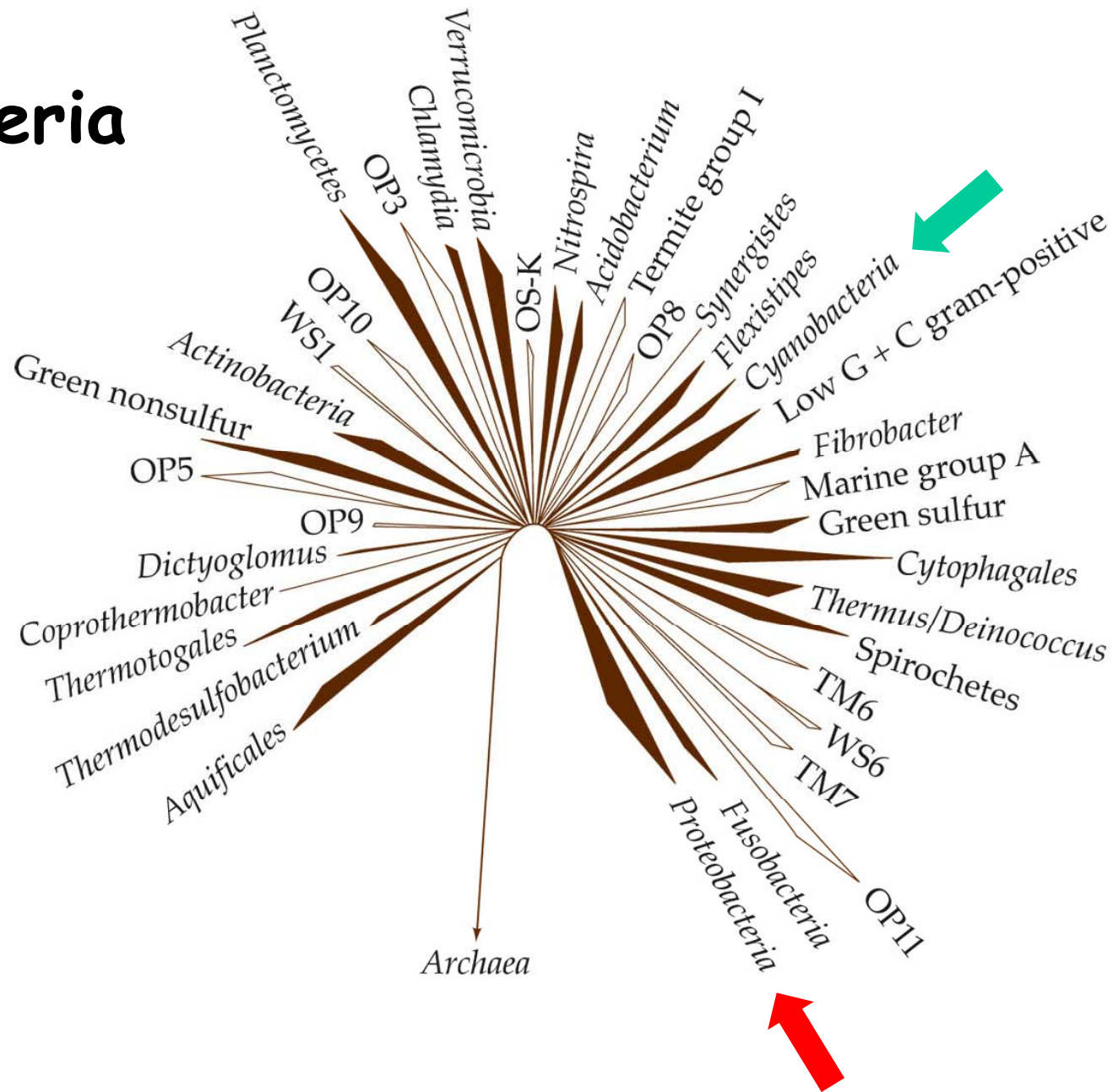
Single origin for all life on Earth...

- 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.

Bacteria



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

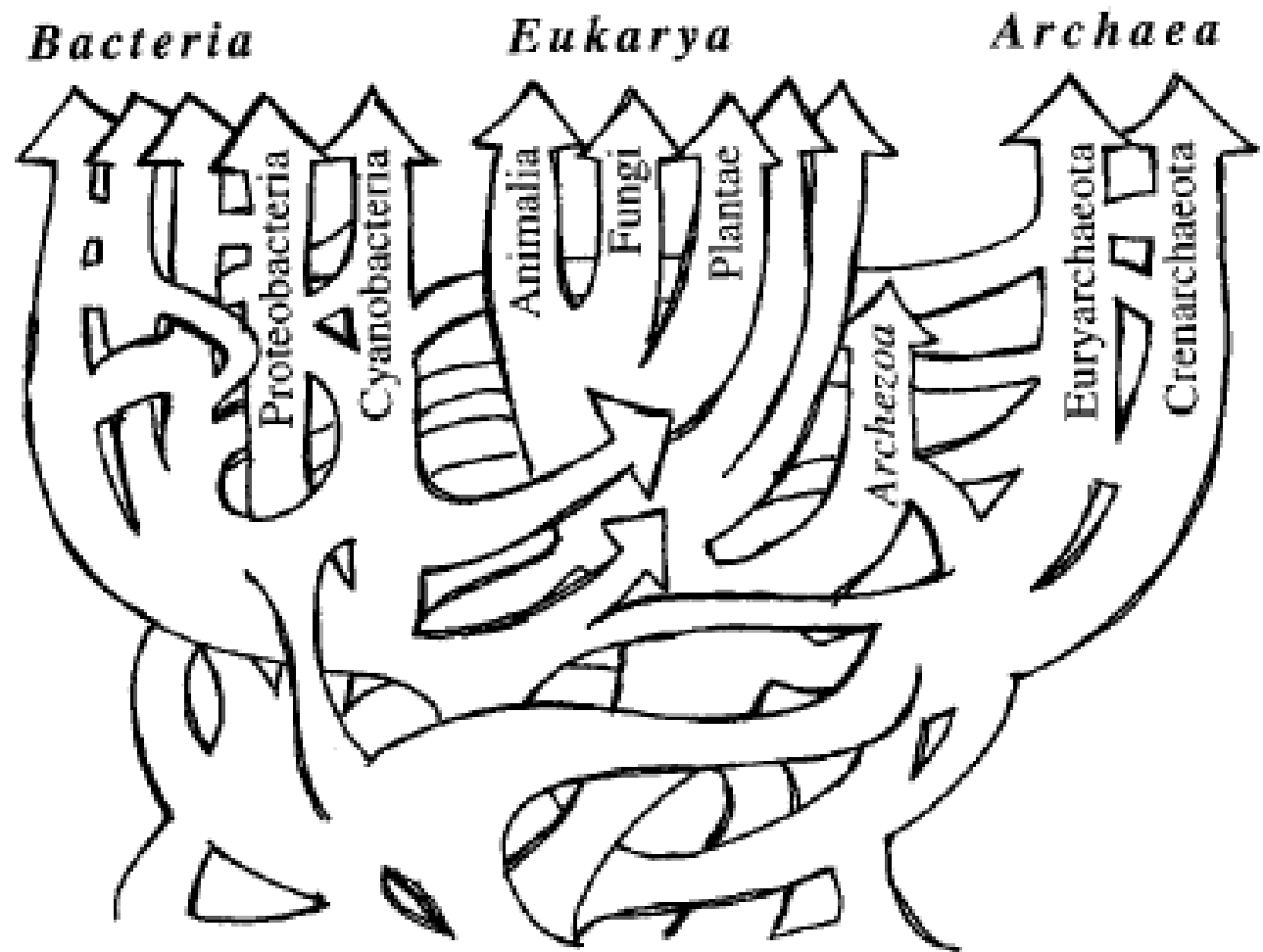
Horizontal gene transfer

This lateral flow of information across microbial taxa occurs via the transfer of genes by:

conjugation, transduction, and transformation.

Rem: These are each one-way processes!

Fig. 3. A reticulated tree, or net, which might more appropriately represent life's history.



Doolittle's Universal Tree (1999)

Gene flow model for two diverging species

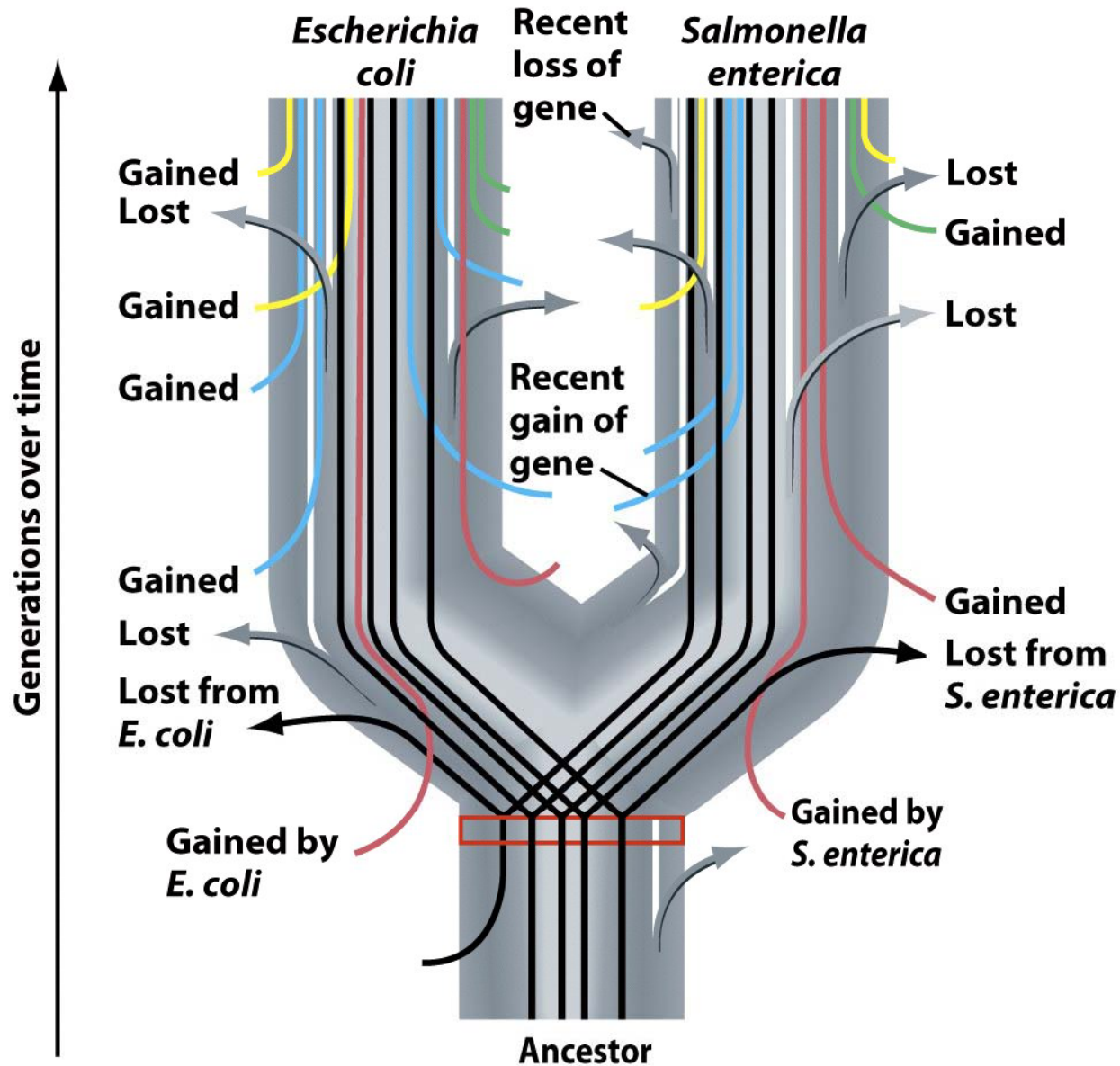
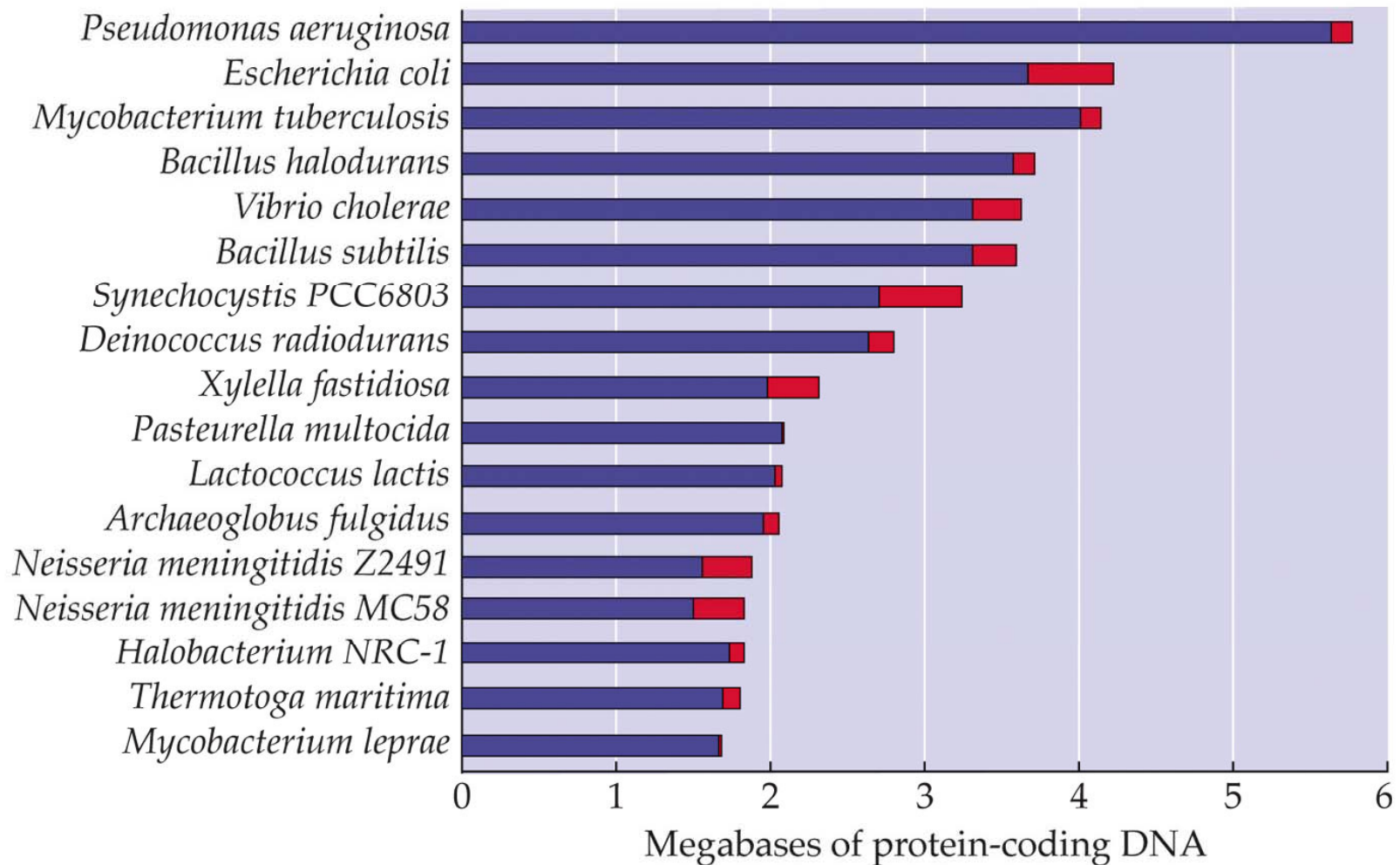
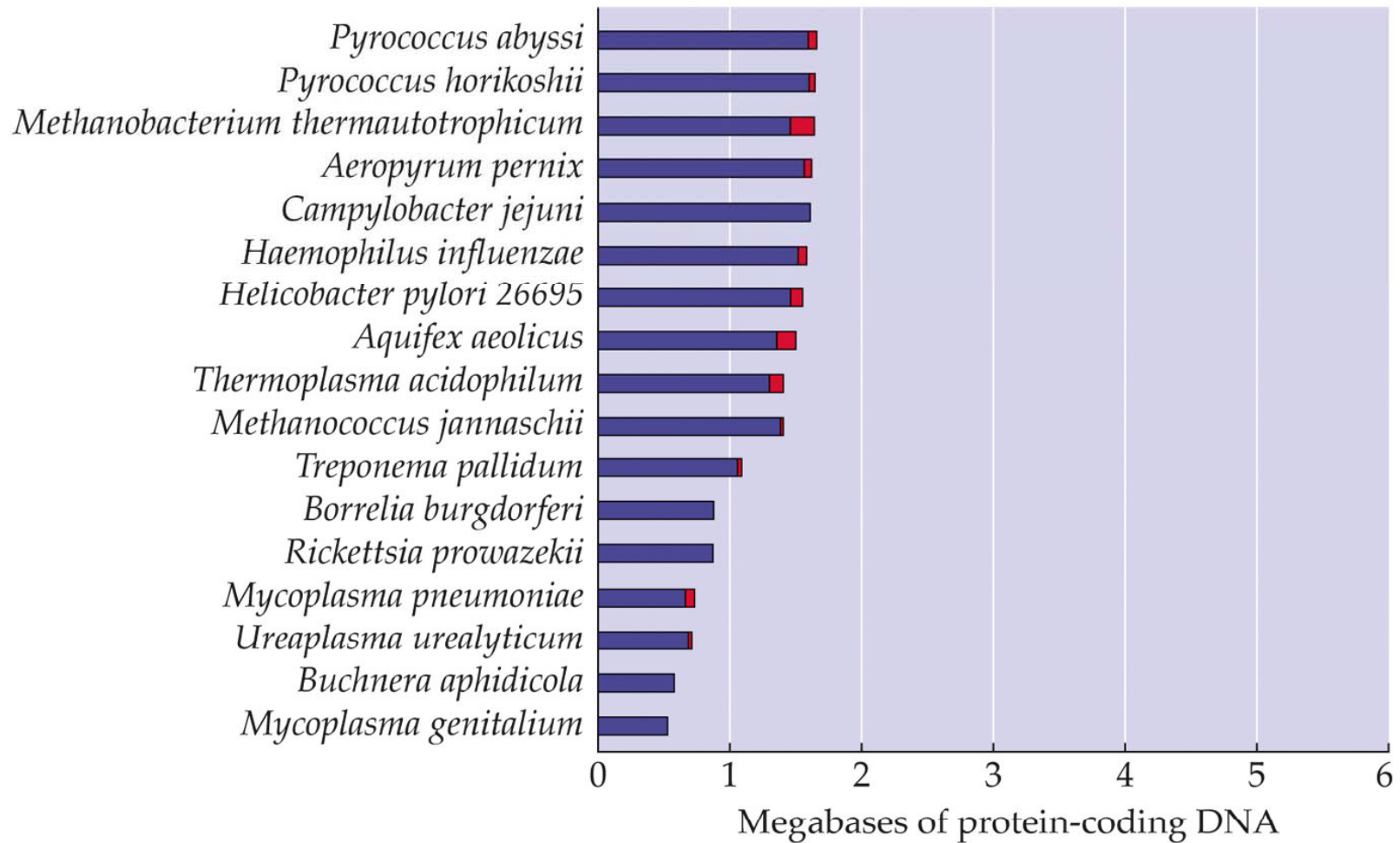


Figure 17.28 Microbiology: An Evolving Science
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Relative importance of horizontal gene transfer



Relative importance of horizontal gene transfer



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.

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** ($\sim 5 \times 10^6$ bp genome), we are **complex** ($\sim 3 \times 10^9$ 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.

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

TABLE 13.3 C values from eukaryotic organisms ranked by size

<i>Species</i>	<i>C value (kb)</i>
<i>Navicula pelliculosa</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>Erysiphe cichoracearum</i> (fungus)	1,500,000
<i>Cyprinus carpio</i> (carp)	1,700,000
<i>Lampraea planeri</i> (lamprey)	1,900,000
<i>Boa 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
<i>Homo sapiens</i> (human)	3,400,000
<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>Protopterus aethiopicus</i> (lungfish)	140,000,000
<i>Ophioglossum petiolatum</i> (fern)	160,000,000
<i>Amoeba proteus</i> (amoeba)	290,000,000
<i>Amoeba dubia</i> (amoeba)	670,000,000



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.

Table 17.2**Comparison of *E. coli* and its primate host species^a**

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

^aAdapted from J. T. Staley, *ASM News*, 1999.

^bValue for all primates.

^cMouse 18S rRNA differs from humans by 16 bases.

^dComparison between *Homo sapiens* and chimpanzee.

^eComparison between *Homo sapiens* and lemurs.

