

*Origin of the Universe – stars, planets, elements*

*Origin of biorelevant monomers – primordial soup*

*Complex chemical processes on the way to living systems*

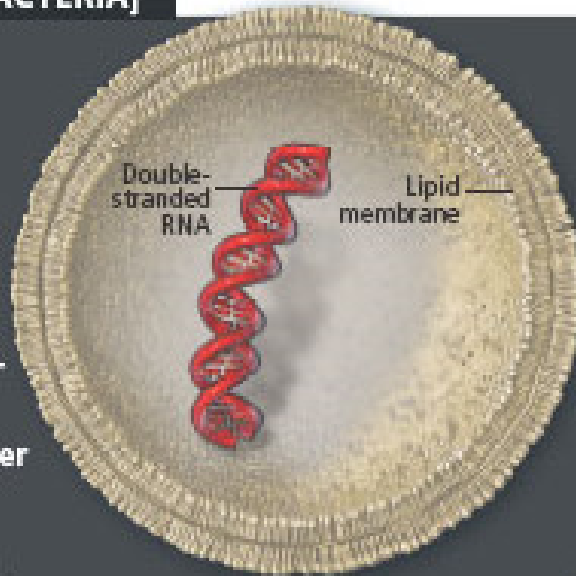
*Protocells and LUCA*

## From RNA world to bacteria

[FROM RNA WORLD TO BACTERIA]

### Journey to the Modern Cell

After life got started, competition among life-forms fueled the drive toward ever more complex organisms. We may never know the exact details of early evolution, but here is a plausible sequence of some of the major events that led from the first protocell to DNA-based cells such as bacteria.

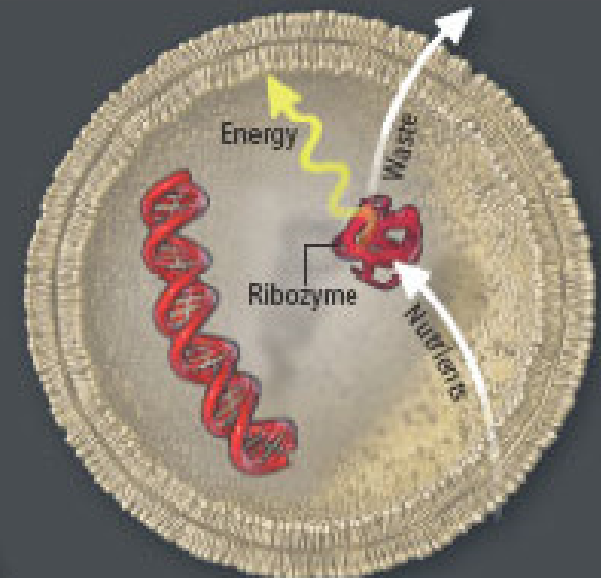
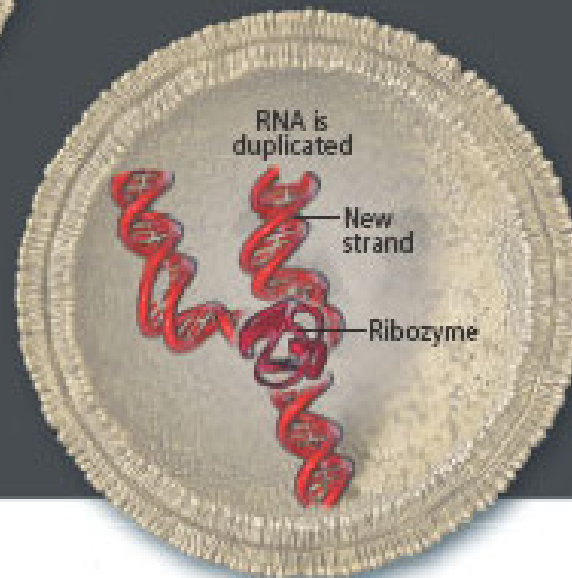


#### 1 EVOLUTION STARTS ▲

The first protocell is just a sac of water and RNA and requires an external stimulus (such as cycles of heat and cold) to reproduce. But it will soon acquire new traits.

#### 2 RNA CATALYSTS ▼

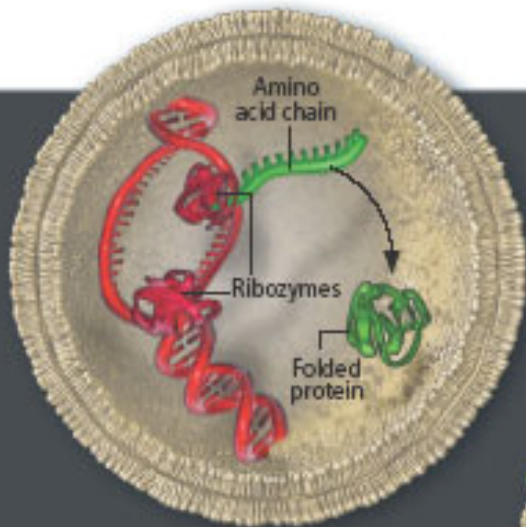
Ribozymes—folded RNA molecules analogous to protein-based enzymes—arise and take on such jobs as speeding up reproduction and strengthening the protocell's membrane. Consequently, protocells begin to reproduce on their own.



#### 3 METABOLISM BEGINS ▲

Other ribozymes catalyze metabolism—chains of chemical reactions that enable protocells to tap into nutrients from the environment.

## From RNA world to bacteria

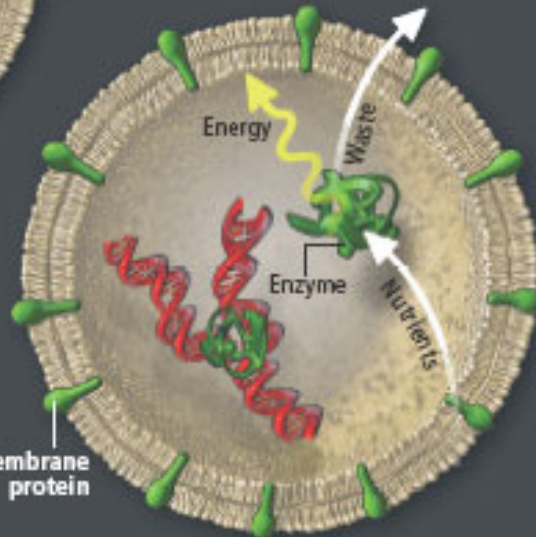


### 4 PROTEINS APPEAR ▲

Complex systems of RNA catalysts begin to translate strings of RNA letters (genes) into chains of amino acids (proteins). Proteins later prove to be more efficient catalysts and able to carry out a variety of tasks.

### 5 PROTEINS TAKE OVER ▼

Proteins take on a wide range of tasks within the cell. Protein-based catalysts, or enzymes, gradually replace most ribozymes.

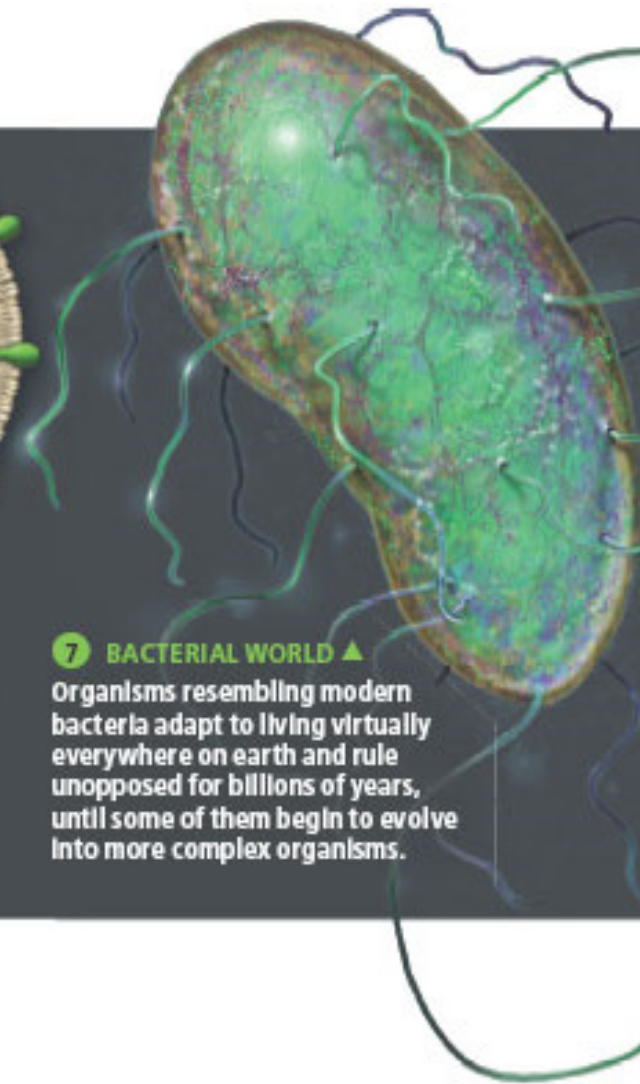


### 6 THE BIRTH OF DNA ▲

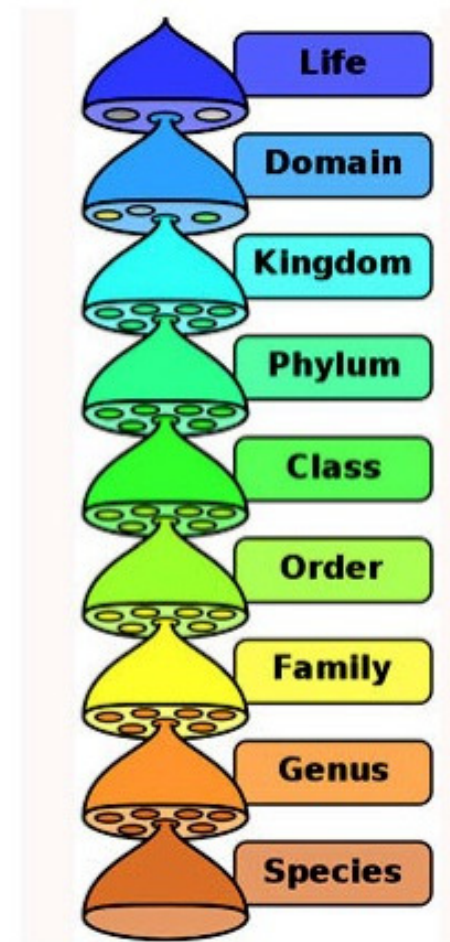
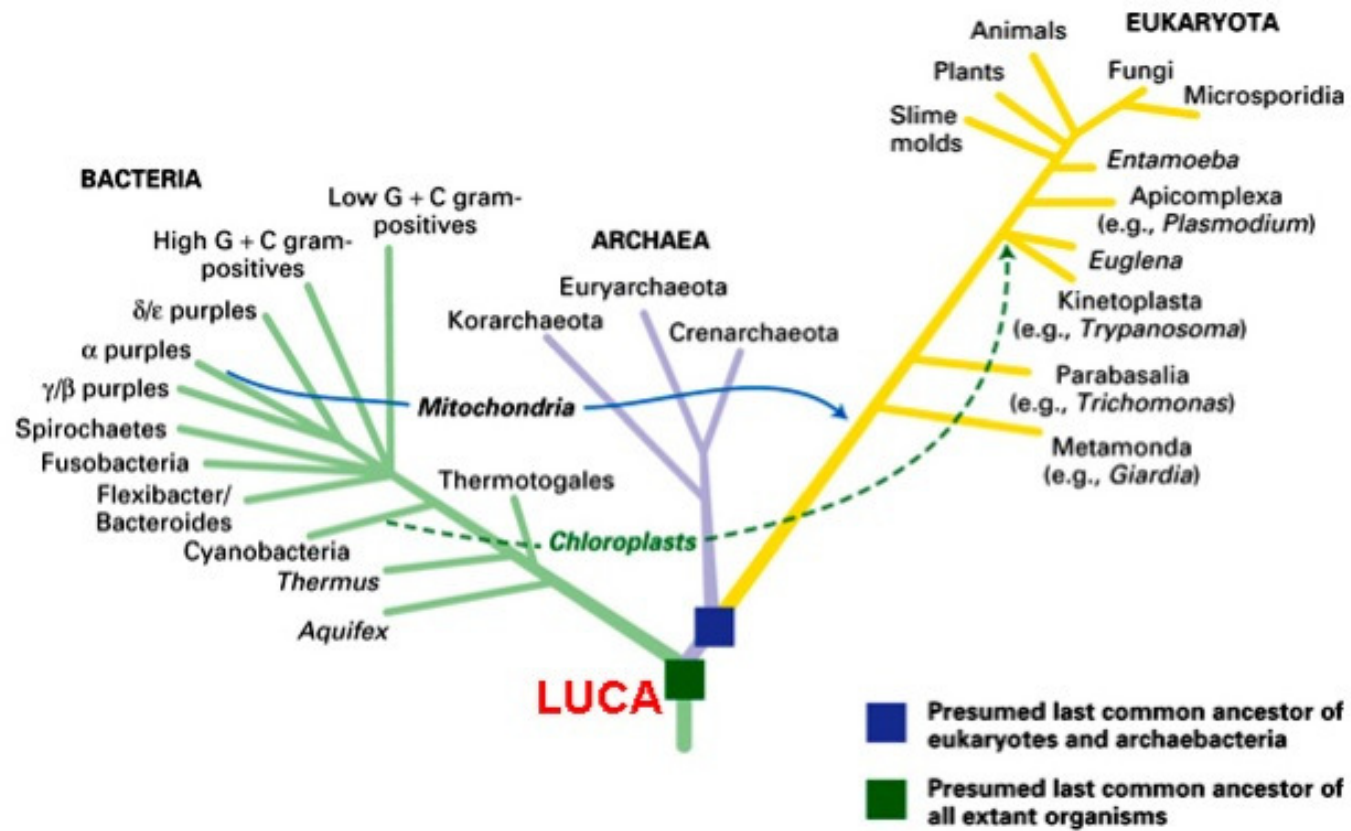
Other enzymes begin to make DNA. Thanks to its superior stability, DNA takes on the role of primary genetic molecule. RNA's main role is now to act as a bridge between DNA and proteins.

### 7 BACTERIAL WORLD ▲

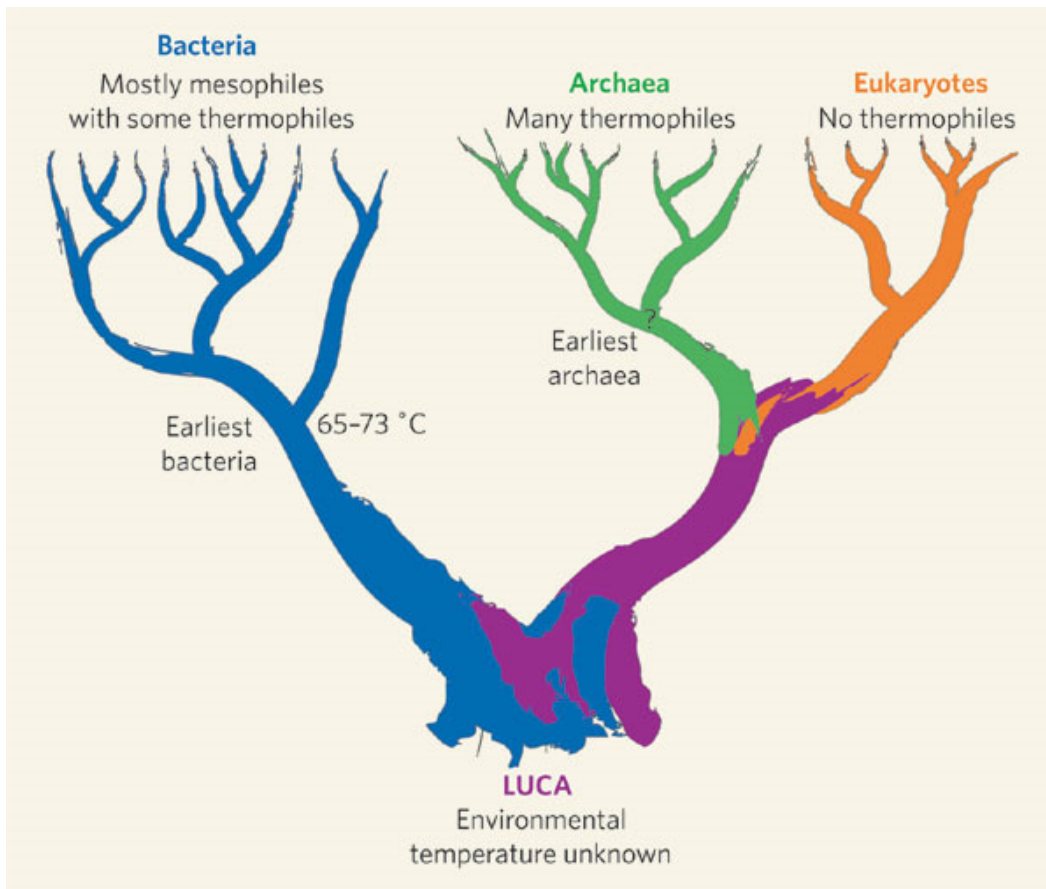
Organisms resembling modern bacteria adapt to living virtually everywhere on earth and rule unopposed for billions of years, until some of them begin to evolve into more complex organisms.







**Figure 1.1a**  
*Molecular Cell Biology*, Seventh Edition  
 © 2013 W.H. Freeman and Company



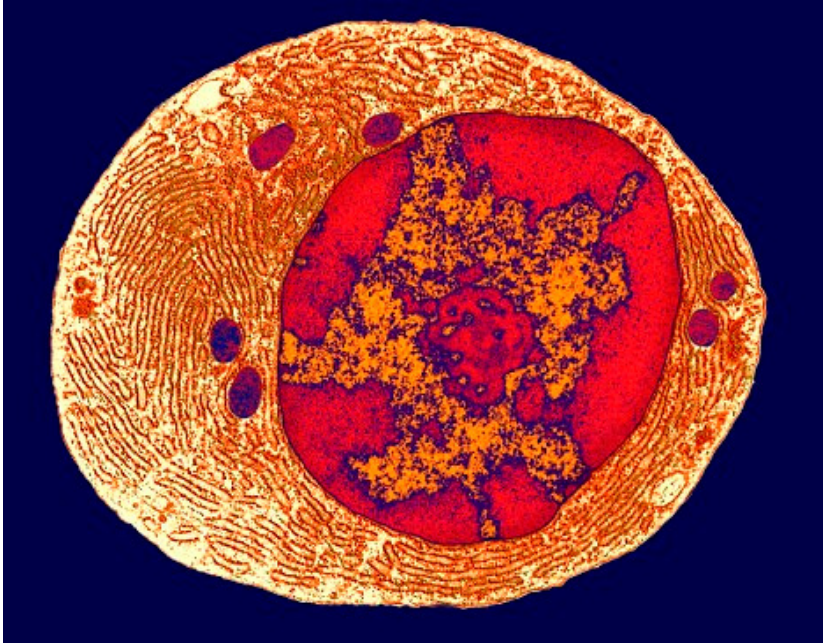
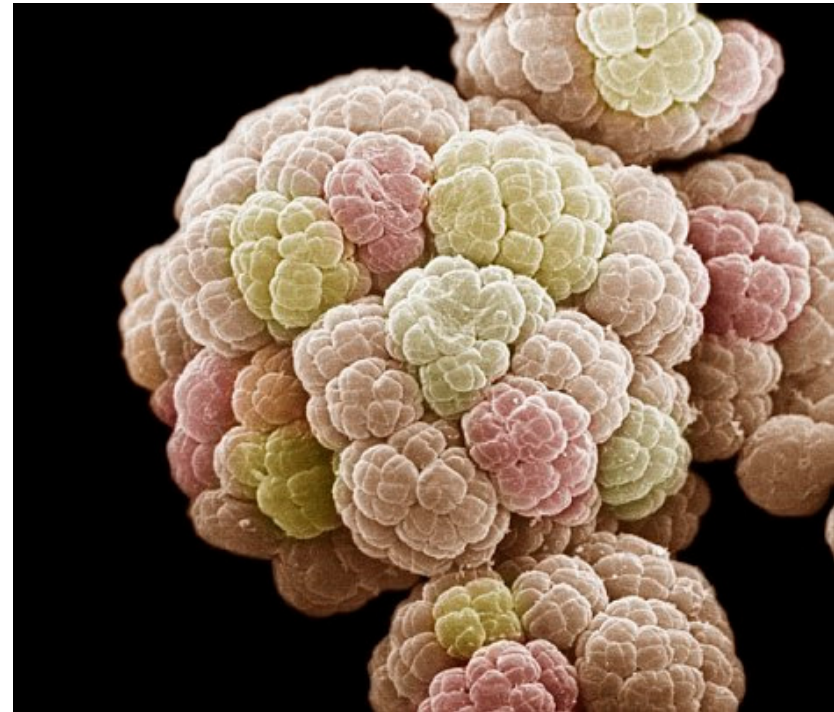


Image of a eukaryotic cell contains numerous organelles, which are now thought to be present in the last universal common ancestor



A colony of the archaea, which form one of the three lines of the tree of life in evolutionary history

# Tree of Life

Billions of Years Ago

Bacteria

Eukarya

Archaea



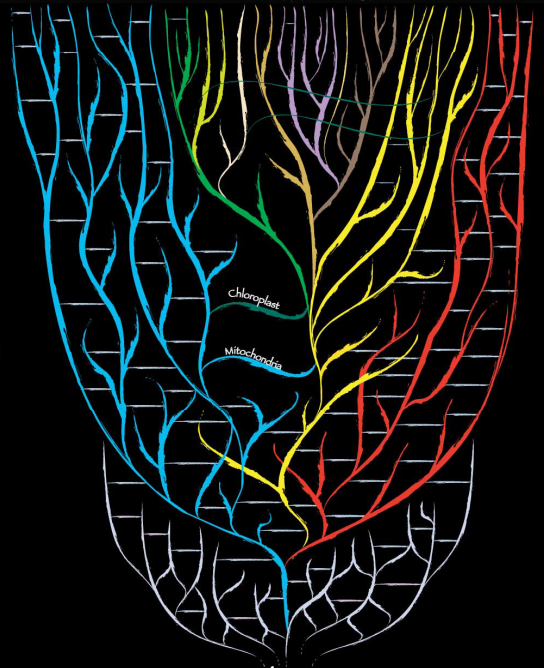
now

1 bya

2 bya

3 bya

4 bya



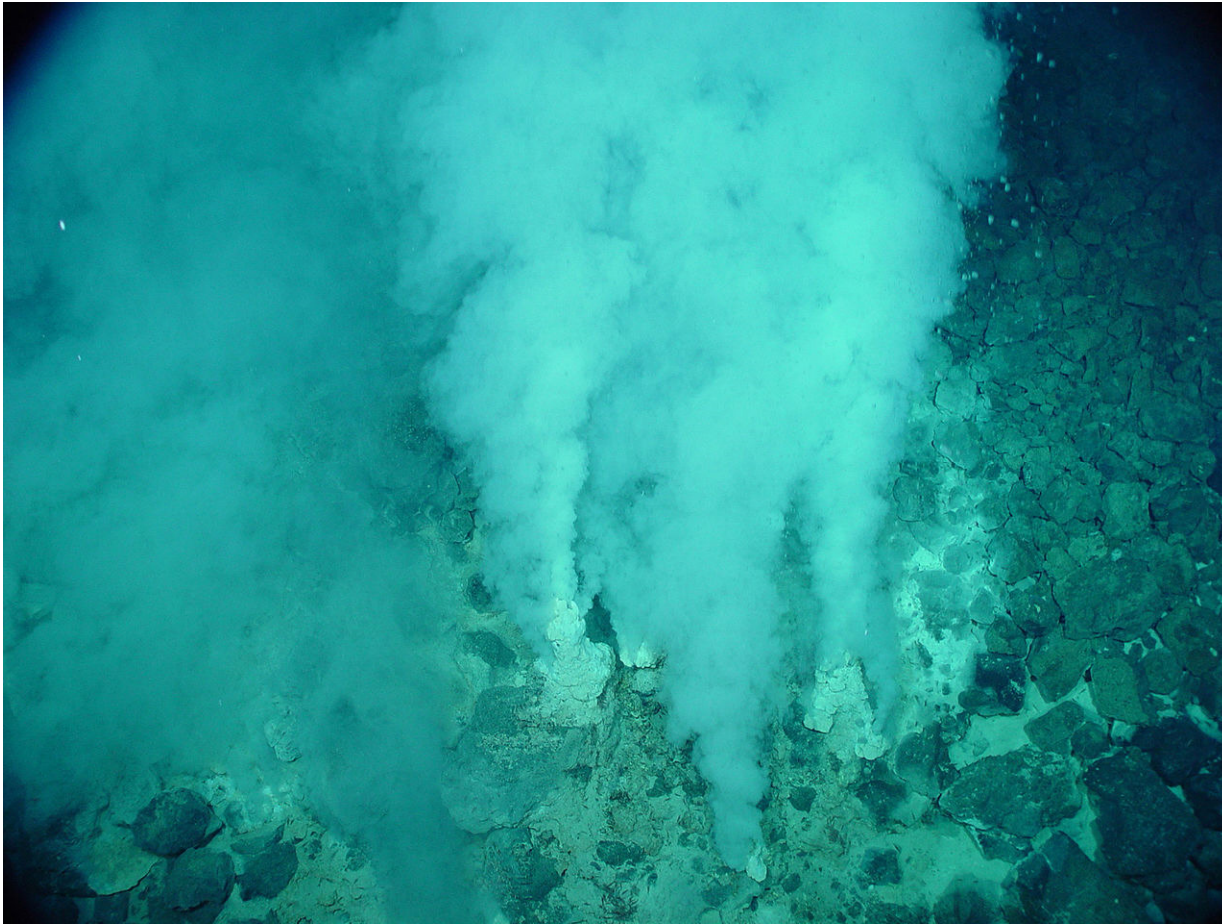
Last Universal Common Ancestor



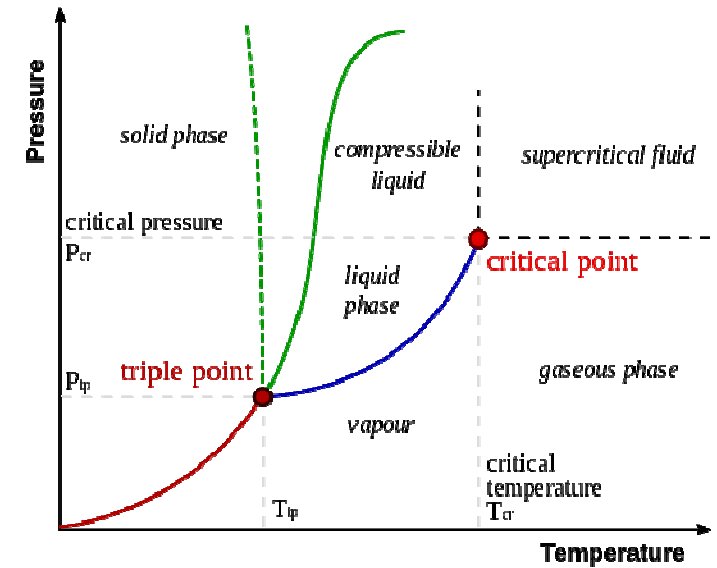


*The Beginning or Origin of Life near Deep Sea Hydrothermal vents*

## Hydrothermal vents

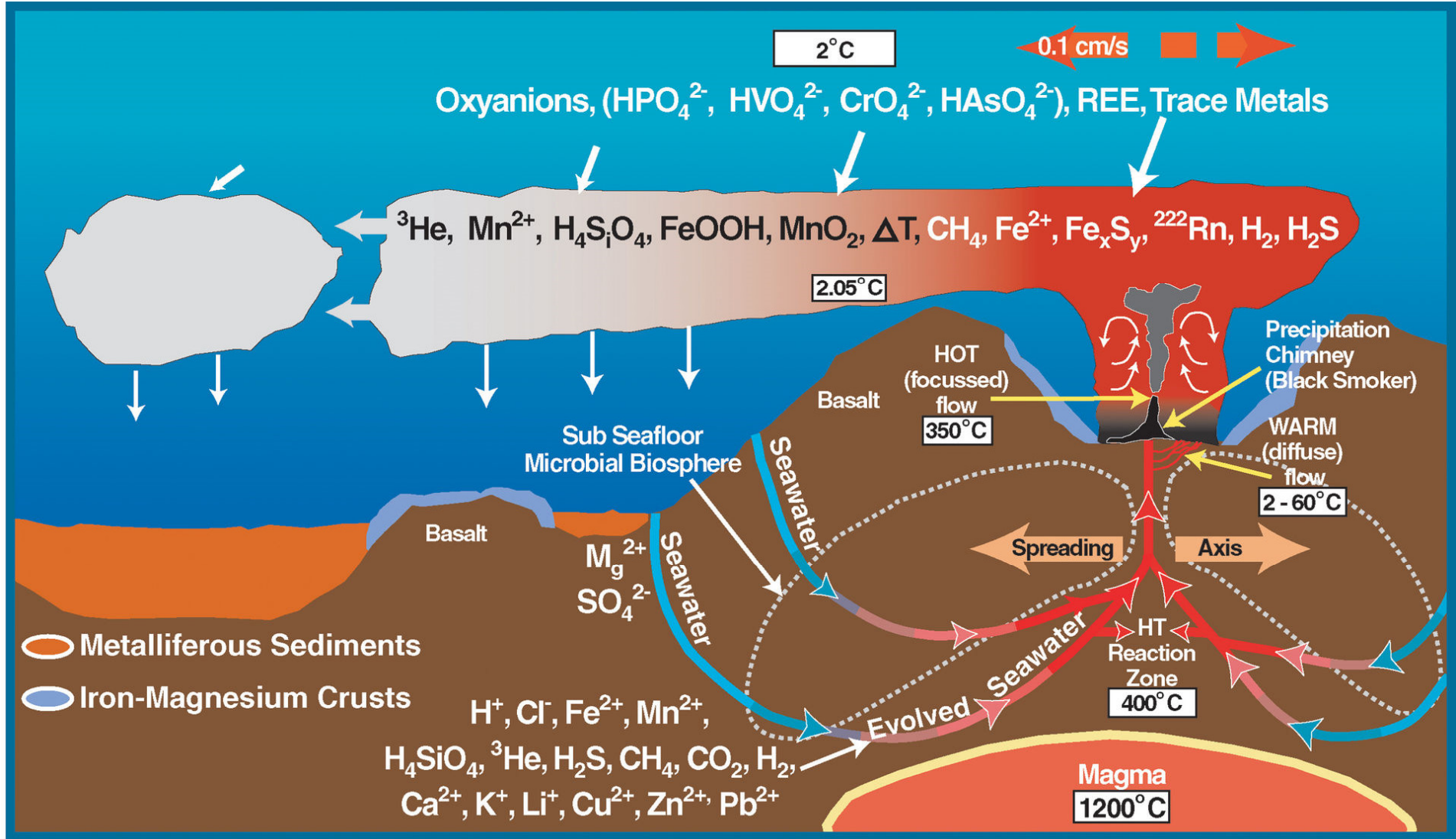


White flocculent mats in and around the extremely gassy, high-temperature ( $>100^{\circ}\text{C}$ ,  $212^{\circ}\text{F}$ ) white smokers at Champagne Vent.

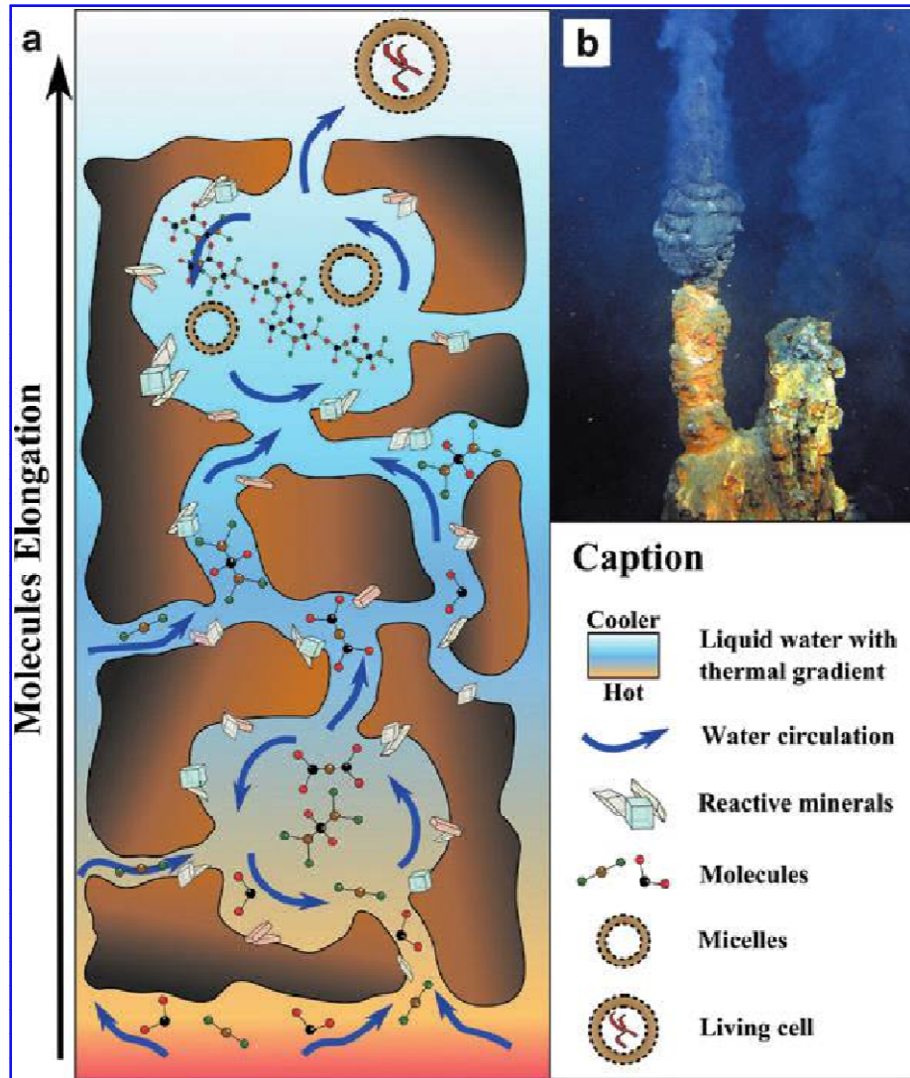


Alkaline hydrothermal vents consist of microscale caverns coated by thin membraneous metal sulfide walls  $\rightarrow$  'Iron-sulfur world'

# Deep sea vent biogeochemical cycle diagram



## Deep sea vent origin of life



### *Possible origin of life in porous hydrothermal vents.*

(a) Sketch showing a porous beehive structure where hydrothermal fluids and seawater can circulate, leading to the accumulation of organic molecules. The reduced mineral surfaces within the vent pores could be favorable locations for the structural organization of macromolecules. We hypothesize the formation of lipid micelles in these environments and the incorporation of information-transferring molecules within the micelles, perhaps due to moderate agitation of the hydrothermal effluent.

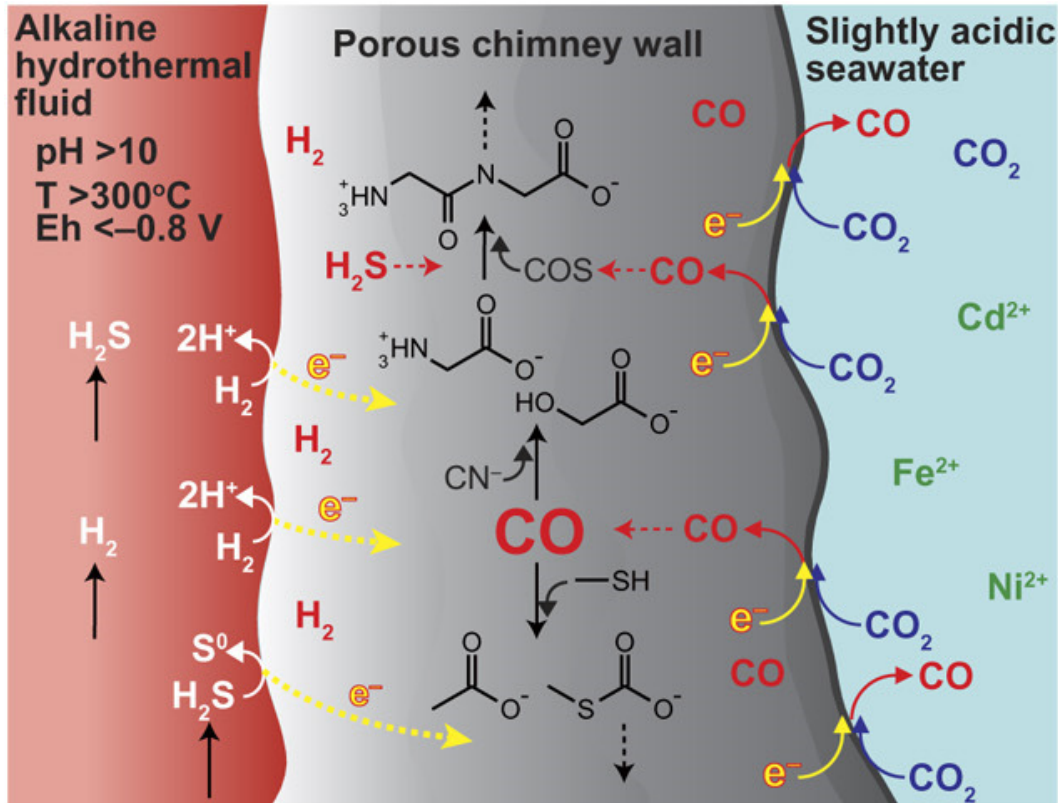
(b) Image of a modern black smoker

(image credit: National Oceanographic and Atmospheric Administration). Color images available online at [www.liebertonline.com/ast](http://www.liebertonline.com/ast)

F. Westall et al., *Astrobiology* **2013**, *13*(9), 887-897



## Deep sea vent origin of life

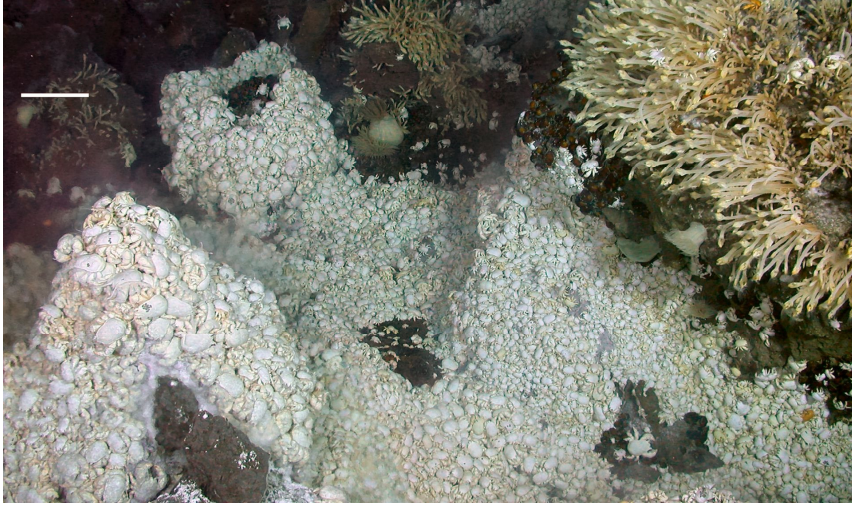


### Abiotic carbon fixation in the primitive hydrothermal system.

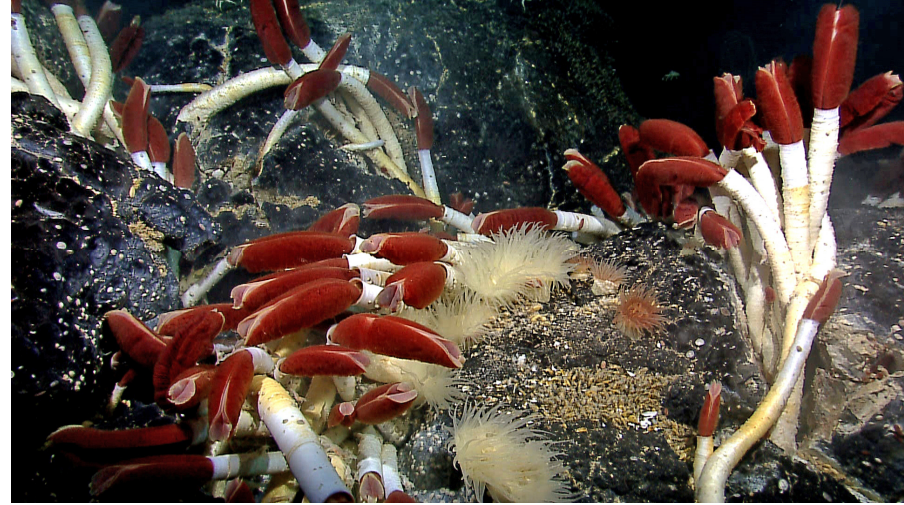
On the ocean floor, mixing of the hydrothermal fluids and seawater generated sulfide-rich chimneys, and the potential gradient across the chimney drove a continuous electron flow. The electric potential at the chimney-seawater interface could reach less than  $-1$  V (versus SHE) in alkaline hydrothermal vent environments. The low potential, in the presence of sulfides rich in  $Cd^{2+}$  and  $Ag^+$ , allowed the electrochemical  $CO_2$  reduction to CO with the FE as high as dozens of percent, together with  $H_2$  evolution. The produced CO served as a driving force for the subsequent abiotic organic synthesis that preceded the origin of life as schematically indicated in the figure

Kitadai et al., *Sci. Adv.* **2018**; 4: eaao7265

## Deep sea vent fauna

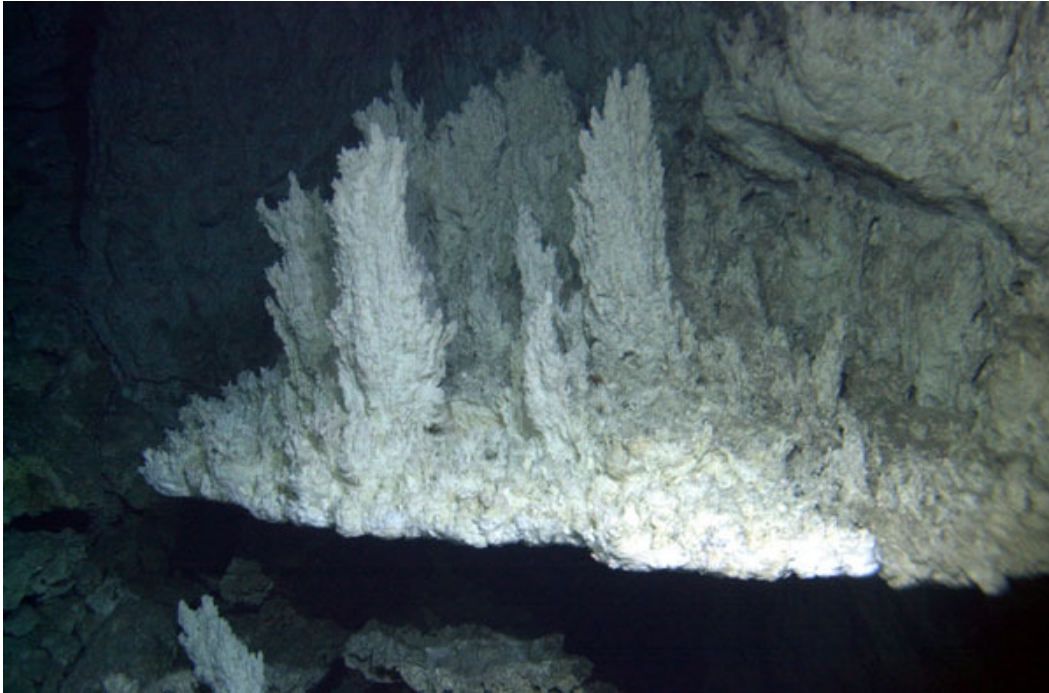


A dense fauna (*Kiwa anomurans* and *Vulcanolepas* like stalked barnacles) near East Scotia Ridge vents



Giant tube worms (*Riftia pachyptila*) cluster around vents in the Galapagos Rift

## *„Lost city” – white smokers: alkaline hydrothermal vents*

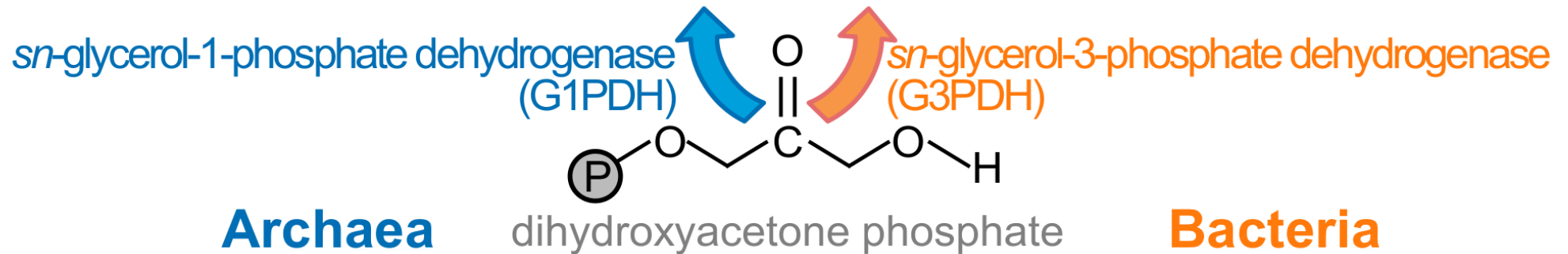
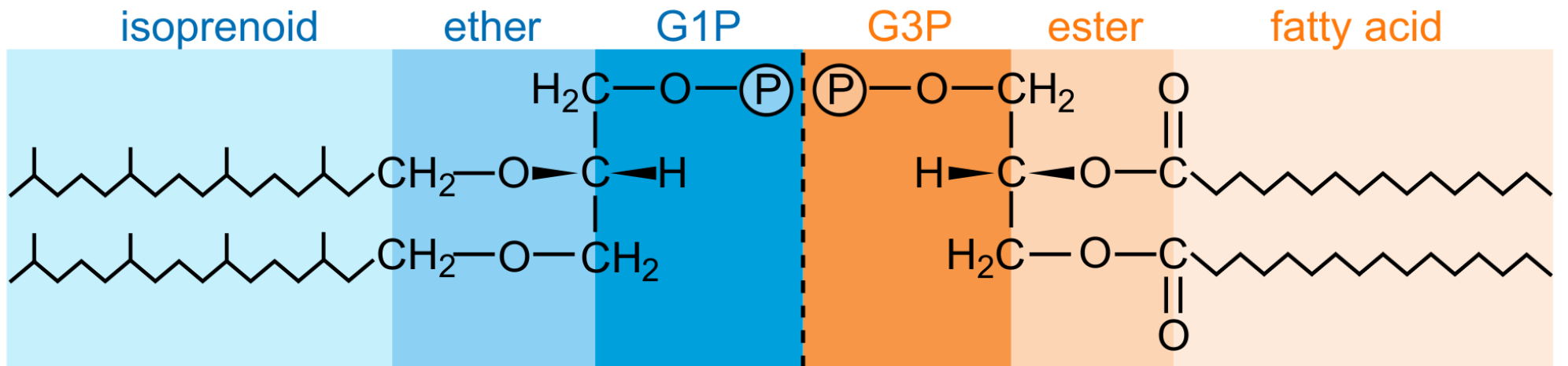


A 1.5-meter-wide ledge on the side of a chimney is topped with dendritic carbonate growths that form when mineral-rich vent fluids seep through the flange and come into contact with the cold seawater.



A carbonate chimney more than 9 meters (30 feet) in height. The white, sinuous spine is freshly deposited carbonate material. The top shows evidence of collapse and re-growth, as indicated by the small newly developed cone on its top

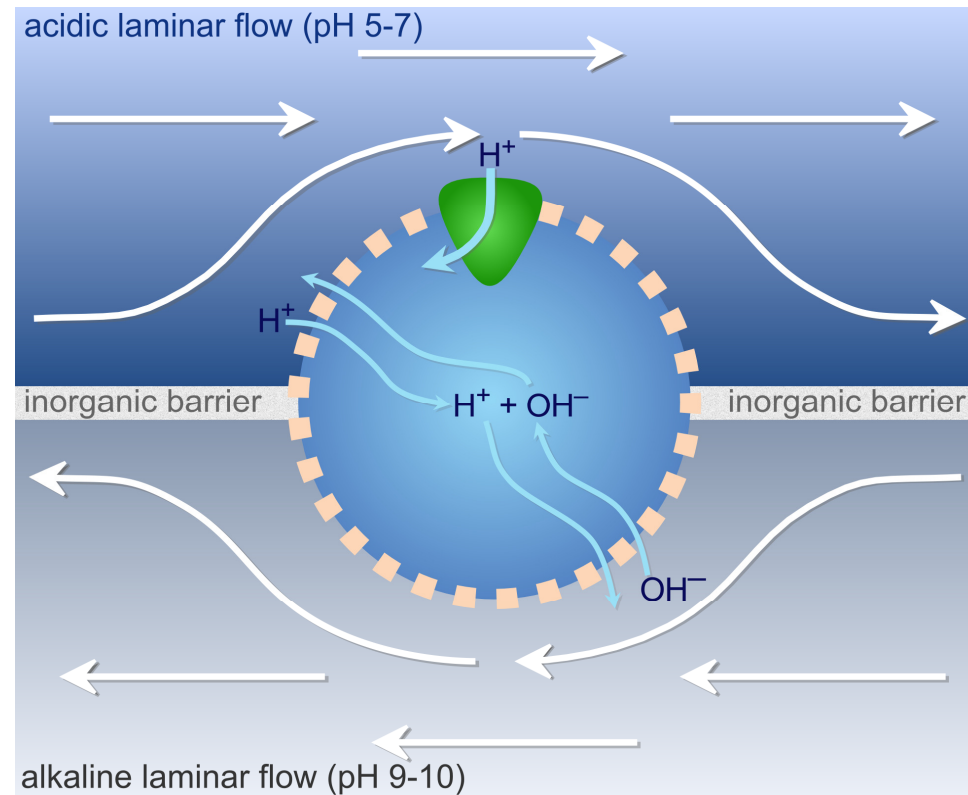




Archaeal lipids: isoprenoid chains + ether bonds + *sn*-glycerol-1-phosphate (G1P) backbone.

Bacterial lipids: fatty acids + ester linkage + *sn*-glycerol-3-phosphate (G3P) skeleton.

Despite widespread horizontal gene transfer, no bacterium has been observed with the archaeal enantiomer, or vice versa. (ether linkages have been observed in bacterial membranes and isoprenoids are common to all three domains)



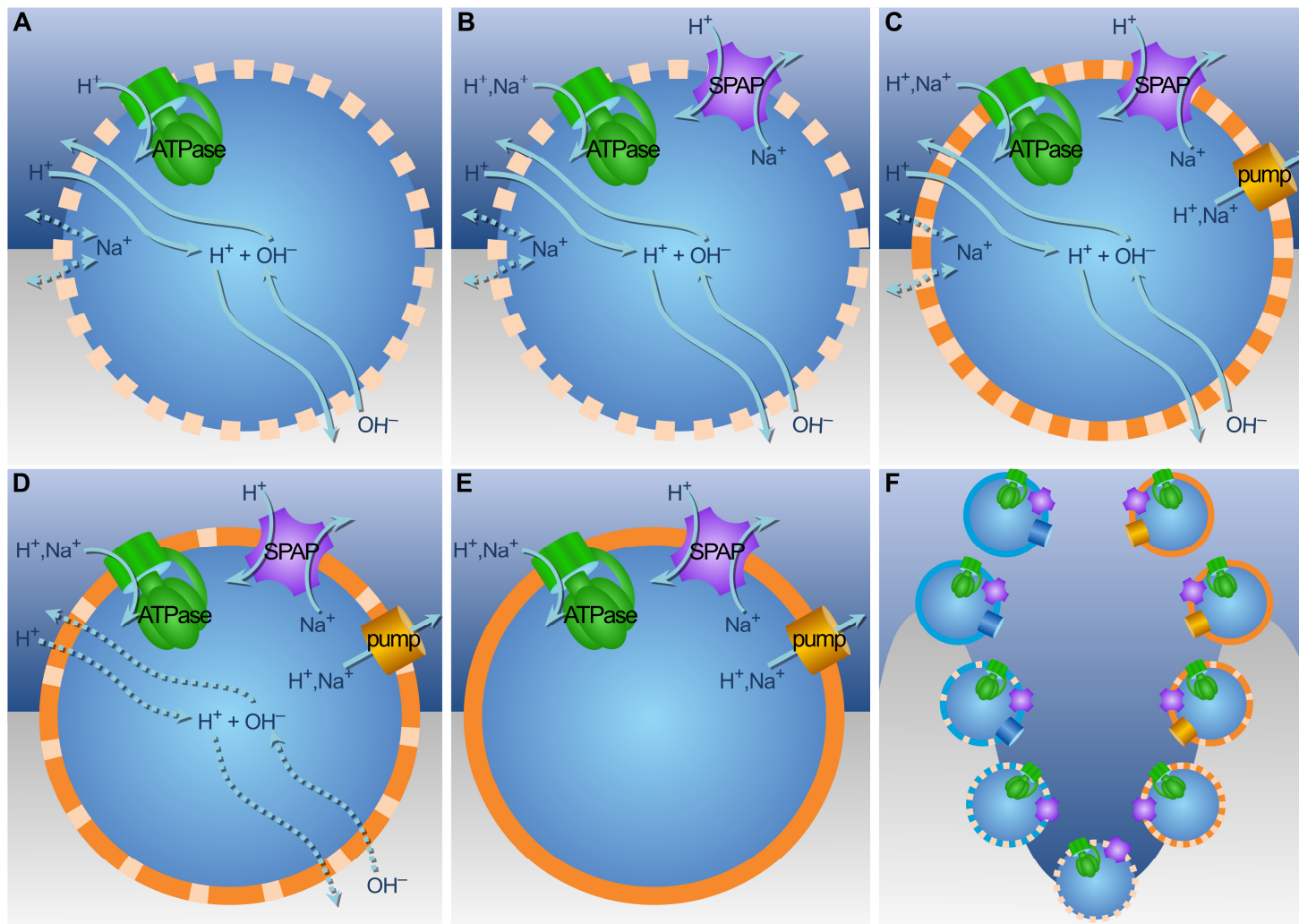
A cell with a semi-permeable membrane at the interface between an alkaline and an acidic fluid (separated elsewhere with an inorganic barrier).  $H^+$ ,  $OH^-$ ,  $Na^+$ ,  $K^+$ ,  $Cl^-$  and other ions flow according to their natural gradients.

Inside the protocell,  $H^+$  and  $OH^-$  can neutralize into water, or leave towards either side.

A protein capable of exploiting the natural proton gradient sits on the acidic side, allowing energy assimilation via ATP production, or carbon assimilation via  $CO_2$  fixation.

V. Sojo, A. Pomiankowski, N. Lane *PLOS Biology*, 2014, 12(8), e1001926

## The role of sodium-proton antiporter (SPAP)



A)  $H^+$  gradient drives energy metabolism (ATPase) or carbon metabolism (Ech)

B) SPAP generates  $Na^+$  from  $H^+$  gradient

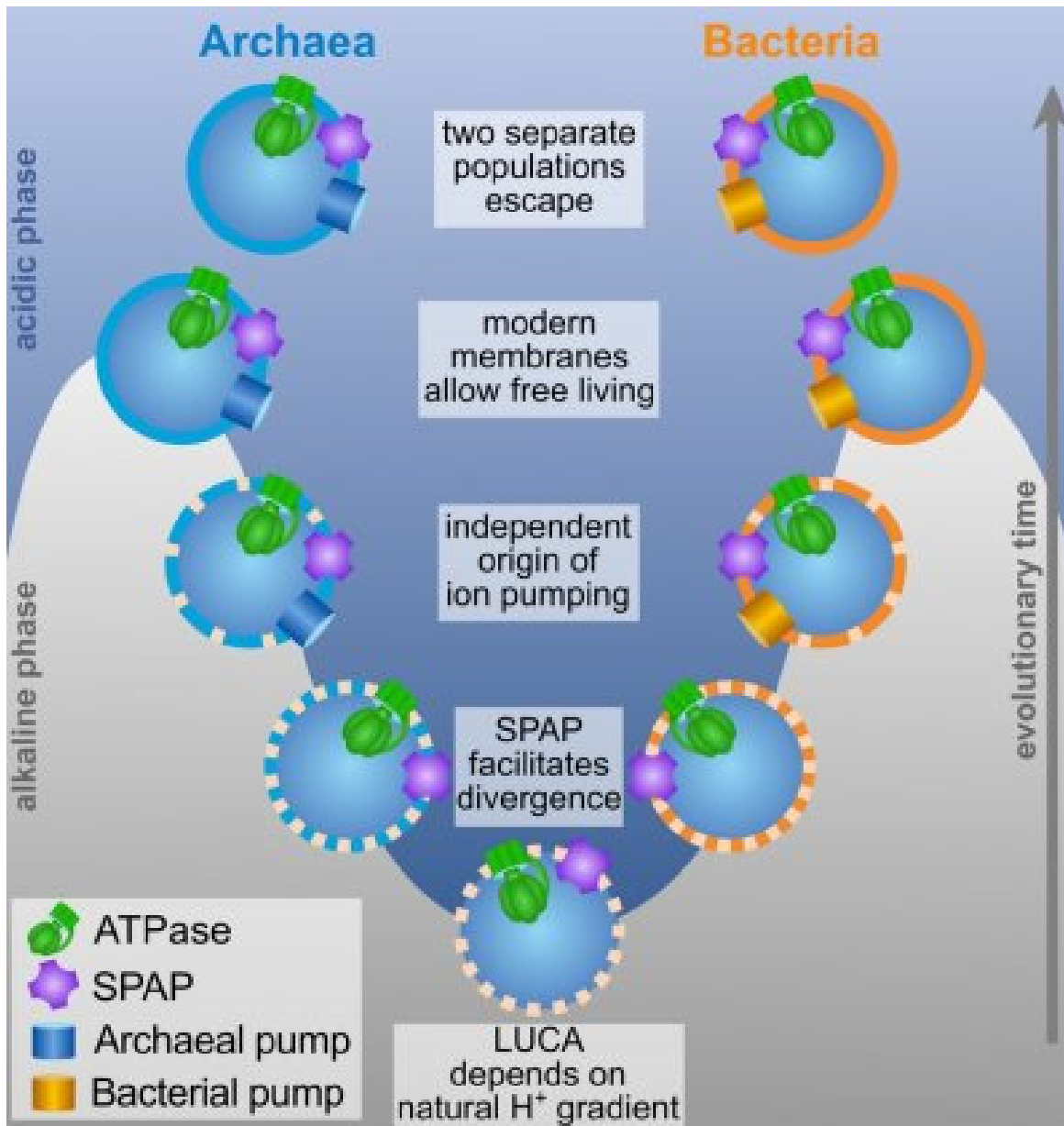
C) Membrane pumps secrete  $H^+$  and  $Na^+$

D) Tighter membranes are now produced, to colonize less alkaline environments

E) Impermeable membranes  $\rightarrow$  gradients created by proteins, independently from the natural environmental gradients

F) SPAP favors divergence, selection for active pumping and tighter membranes; independent evolution of archaea and bacteria

V. Sojo, A. Pomiankowski, N. Lane  
*PLOS Biology*, 2014, 12(8), e1001926



### Origin of autotrophy and development of cell membrane

Ion pumping and phospholipid membranes evolved independently in bacteria and archaea.

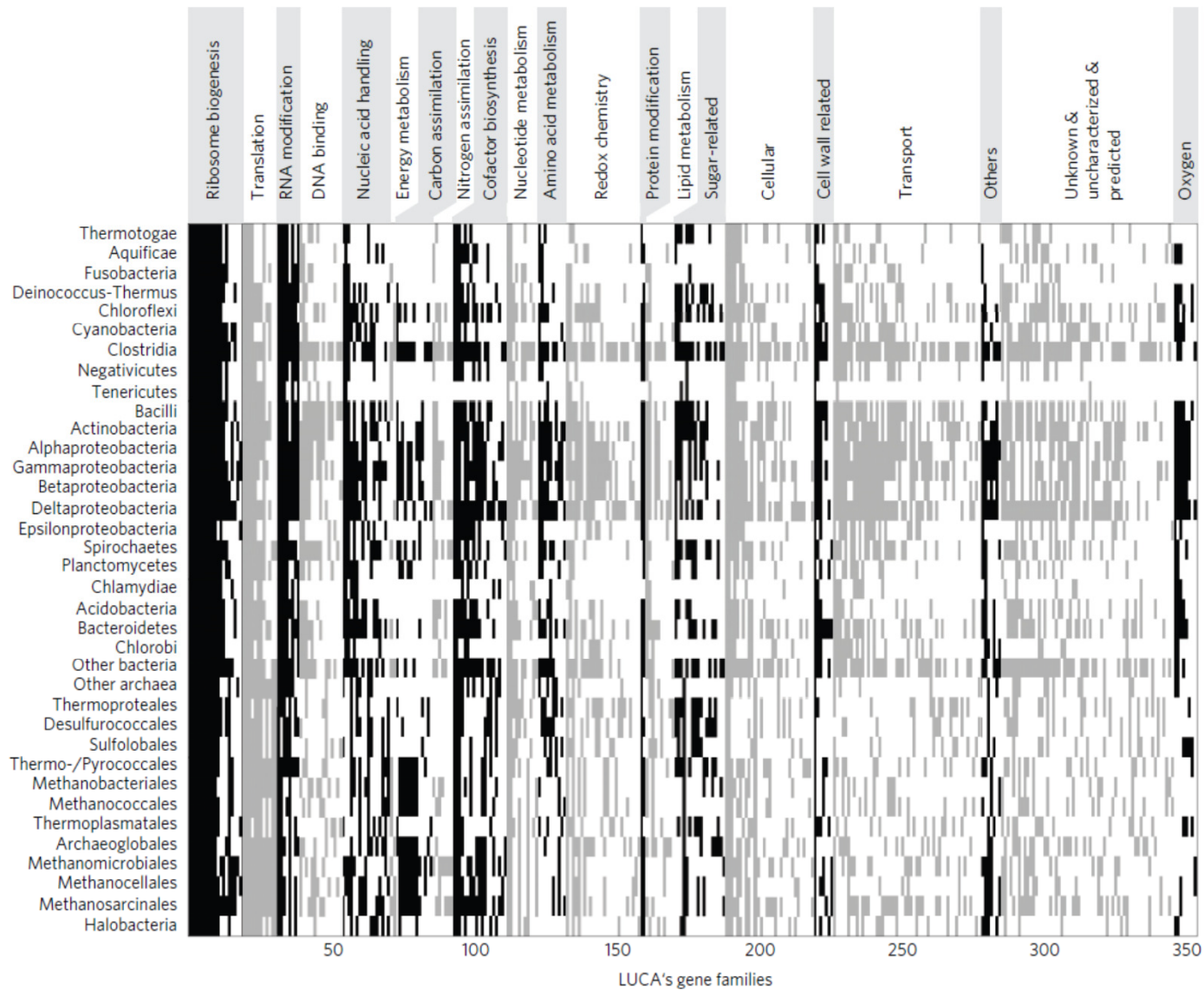
Energy to LUCA could have been delivered by the natural proton gradient in alkaline hydrothermal vents, if the membrane was much more leaky than contemporary ones.

Development of proton pumping allowed for escape from the vent environment.

*sodium-proton antiporter (SPAP)*

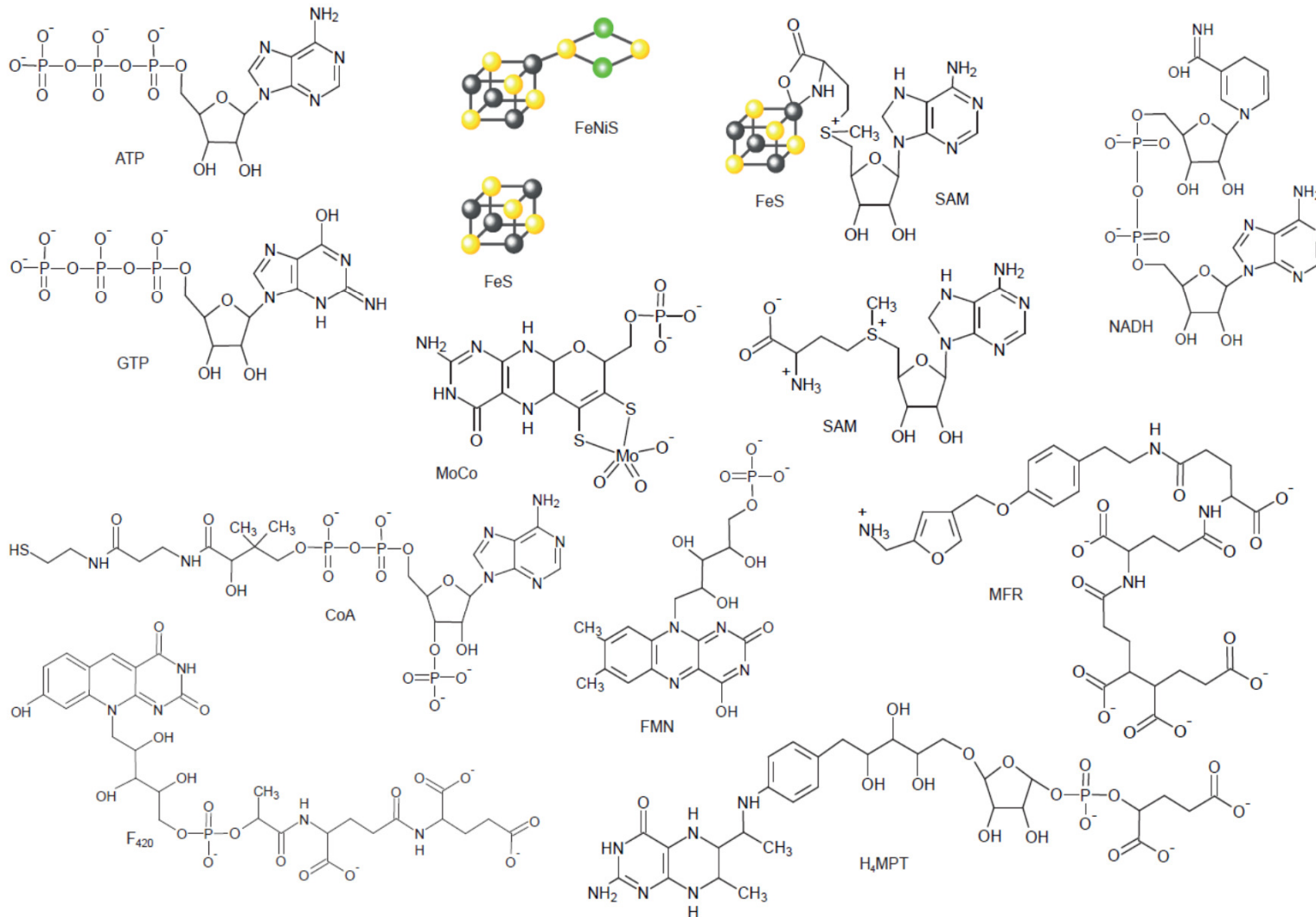
V. Sojo, A. Pomiankowski, N. Lane  
*PLOS Biology*, 2014, 12(8), e1001926

# Taxonomic distribution of LUCA's genes grouped by functional categories



M.C. Weiss et al. *Nature Microbiology*,  
2016, Article 16116

## Structures of the cofactors found in LUCA's protein set.



FeNiS – nickel-iron-sulfur cluster

FeS – iron-sulfur cluster

MoCo – molybdenum cofactor

SAM – S-adenosylmethionine

CoA – coenzyme A

MFR – methanofuran

H4MPT – tetrahydromethanopterin

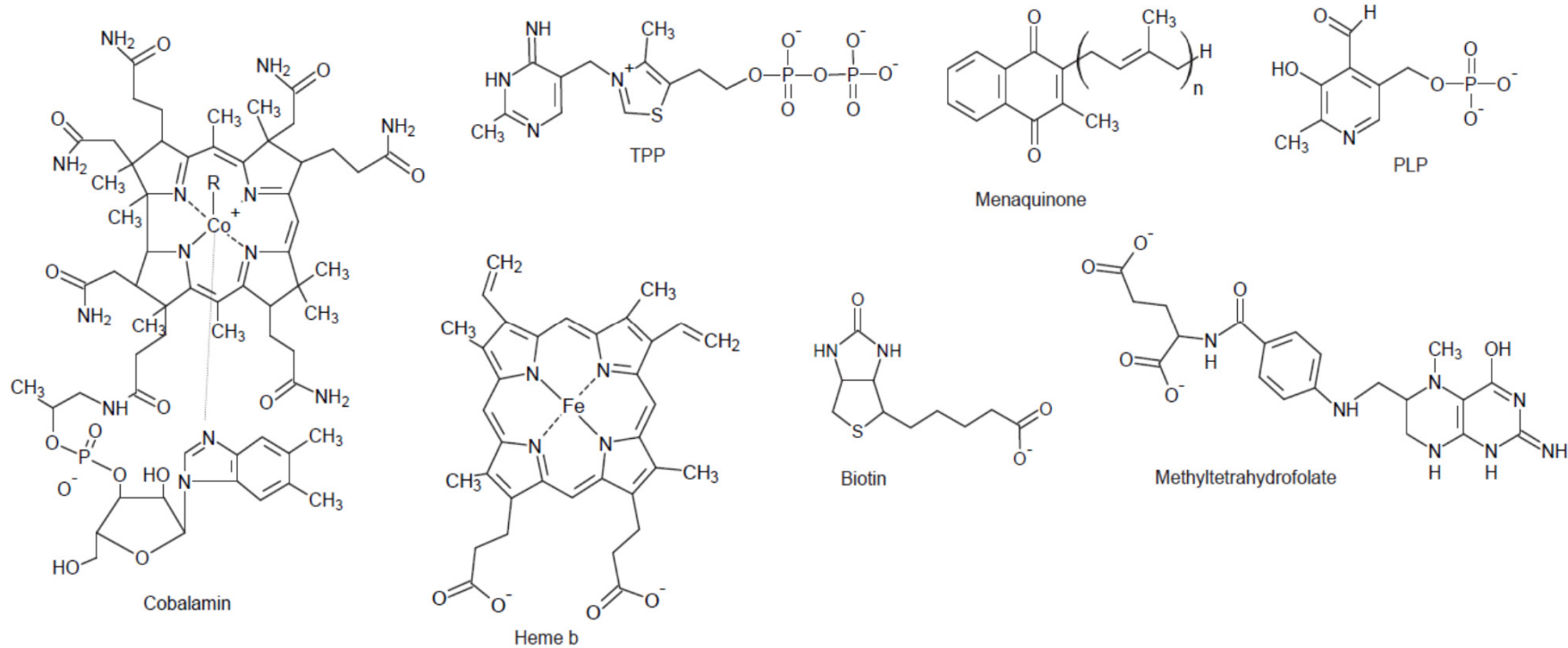
TPP - thiamine pyrophosphate

PLP - pyridoxal phosphate

NTP – nucleoside triphosphate.

M.C. Weiss et al. *Nature Microbiology*,  
2016, Article 16116

## Structures of the cofactors found in LUCA's protein set.



Mononuclear metal centers (Fe and Cu) and the non-standard amino acid selenocysteine are not shown, nor are small protein electron carriers such as ferredoxin or rubredoxin. NTP is also listed as a cofactor, but not shown here as it stands for any of the nucleoside triphosphates in those cases when it's not known which one is bound by the enzyme, or when more than one nucleoside triphosphate can be used

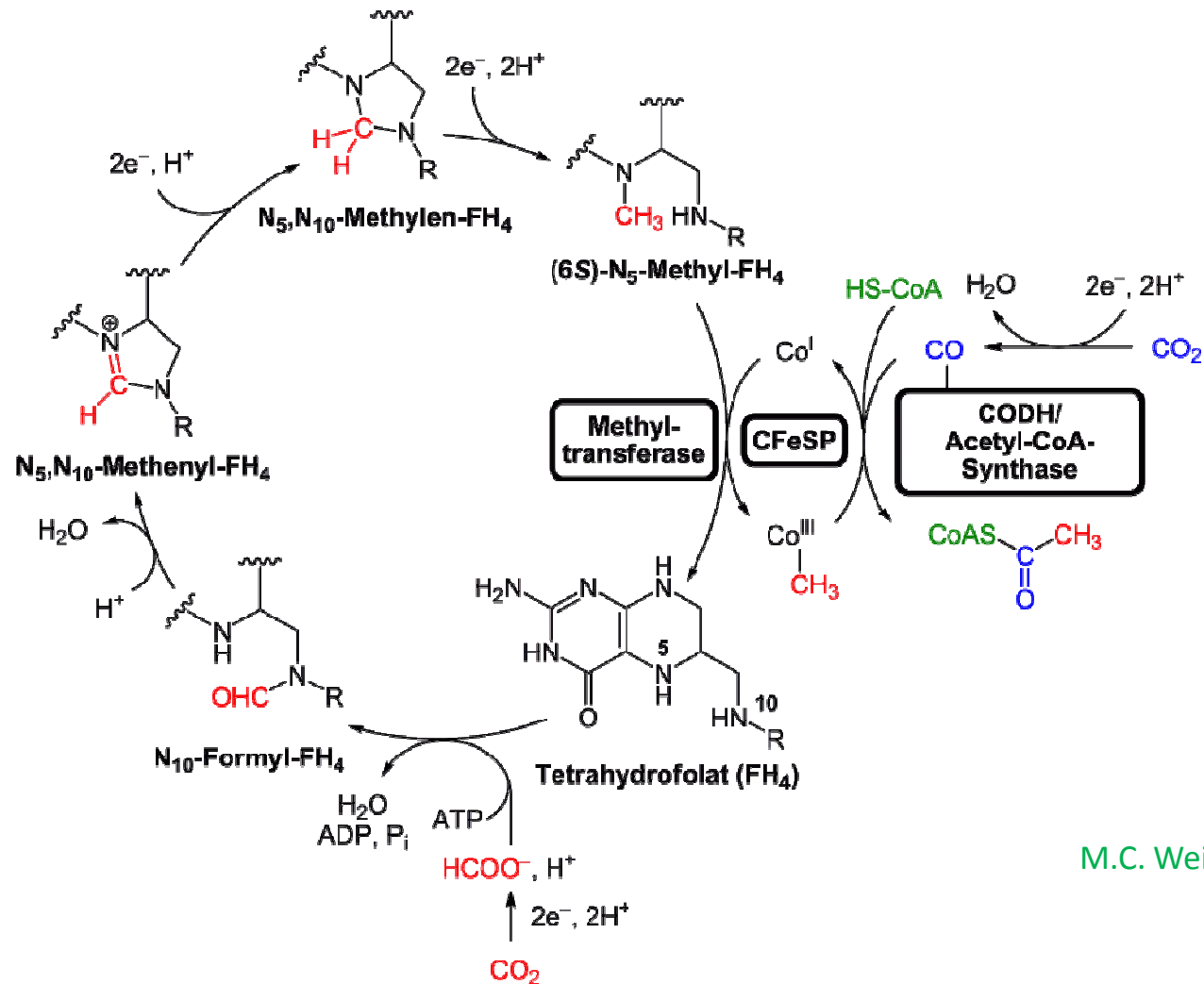
## *Phylogenetic identification of LUCA's proteome*

355 protein families shared among contemporary *archaea* and *bacteria*, including:

- 19 proteins involved in ribosome biogenesis
- 8 aminoacyl tRNA synthetases
- proteins for carbon, energy, and nitrogen metabolism
- rotor-stator ATP synthase subunit (ion gradients were likely supplied geochemically)
- substrate-level phosphorylation (acetylphosphate from acetyl-CoA)
- reverse gyrase – specific for currently living hyperthermophilic organisms
- chemolithoautotrophy enzymes present (WL pathway), chemoorganoautotrophy enzymes absent



## Wood-Ljungdahl (WL) anaerobic pathway of carbon fixation



M.C. Weiss et al. *Nature Microbiology*,  
2016, Article 16116

A primitive metabolic pathway for carbon fixation, still used by some contemporary chemoautotrophic organisms

## *Metabolism of LUCA*

Among six currently known pathways of CO<sub>2</sub> fixation, only WL pathway was present in LUCA:

The relevant enzymes are packed with FeS and FeNiS centres

They require cofactors: flavin, F<sub>420</sub>, methanofuran, two pterins and corrins

Hydrogenases also present in LUCA's genome → electrons likely obtained from hydrogen, as in modern microbes using the WL pathway

Nitrogenase and glutamine synthetase serve for nitrogen fixation

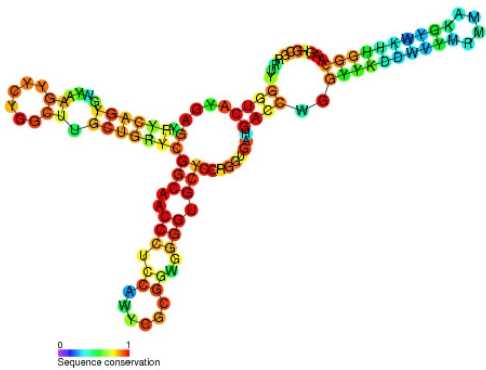
WL pathway, nitrogenase and hydrogenases are very oxygen-sensitive  
→ LUCA was an anaerobic autotroph that could live from gases H<sub>2</sub>, CO<sub>2</sub>, and N<sub>2</sub>.

## Metabolism of LUCA

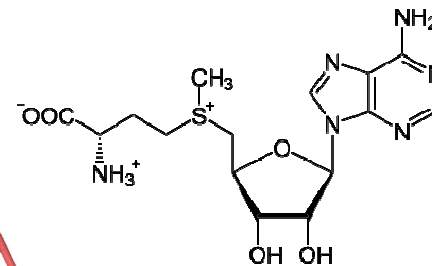
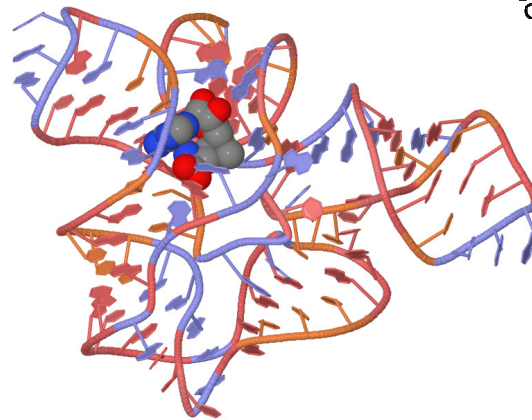
Enzymes for cofactor biosynthesis, including pterins, MoCo, cobalamin, siroheme, TPP, CoM and F<sub>420</sub>, are also conserved.

Many of them are S-adenosyl methionine(SAM)-dependent

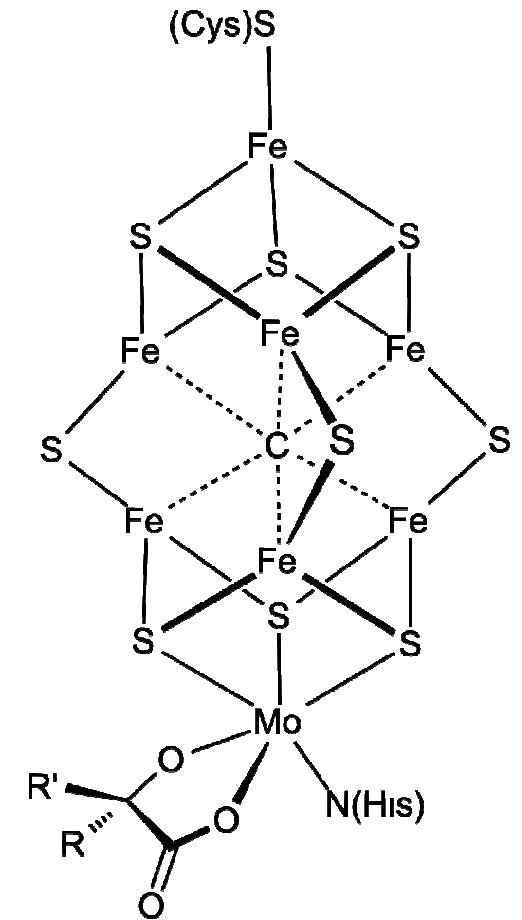
SAM chemistry is based on oxygen-sensitive FeS-containing proteins that initiate radical-dependent methylations.



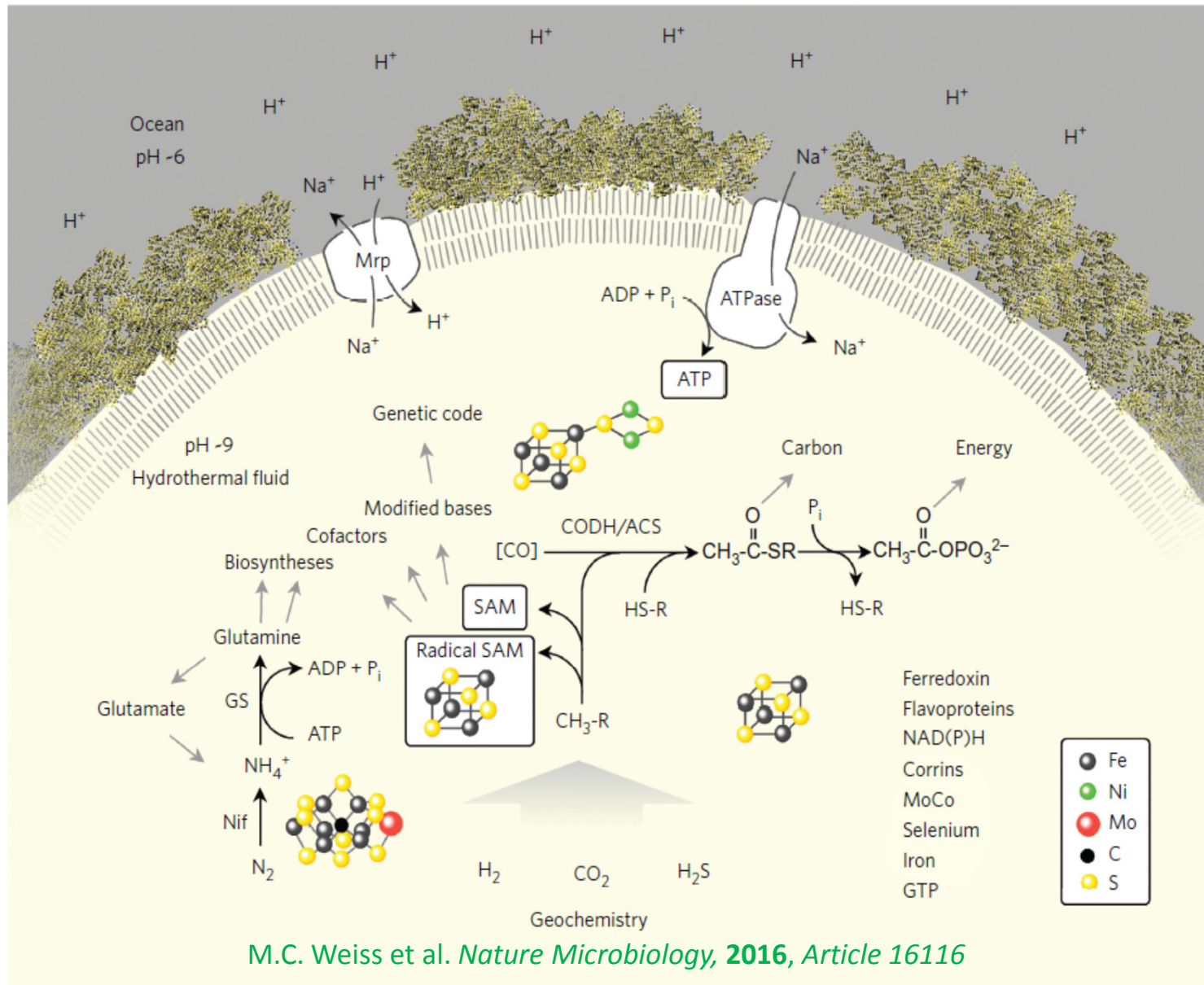
*SAM-binding riboswitches*



M.C. Weiss et al. *Nature Microbiology*,  
2016, Article 16116

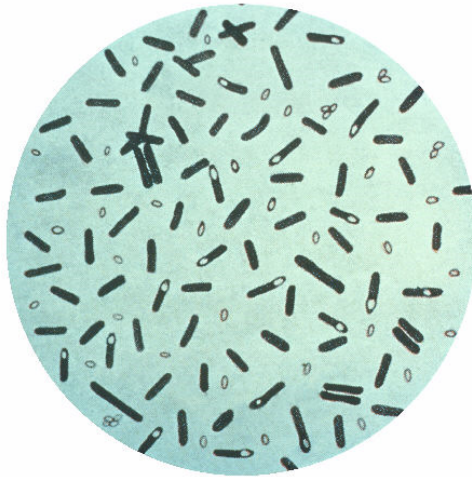


*FeMo cofactor of nitrogenase*

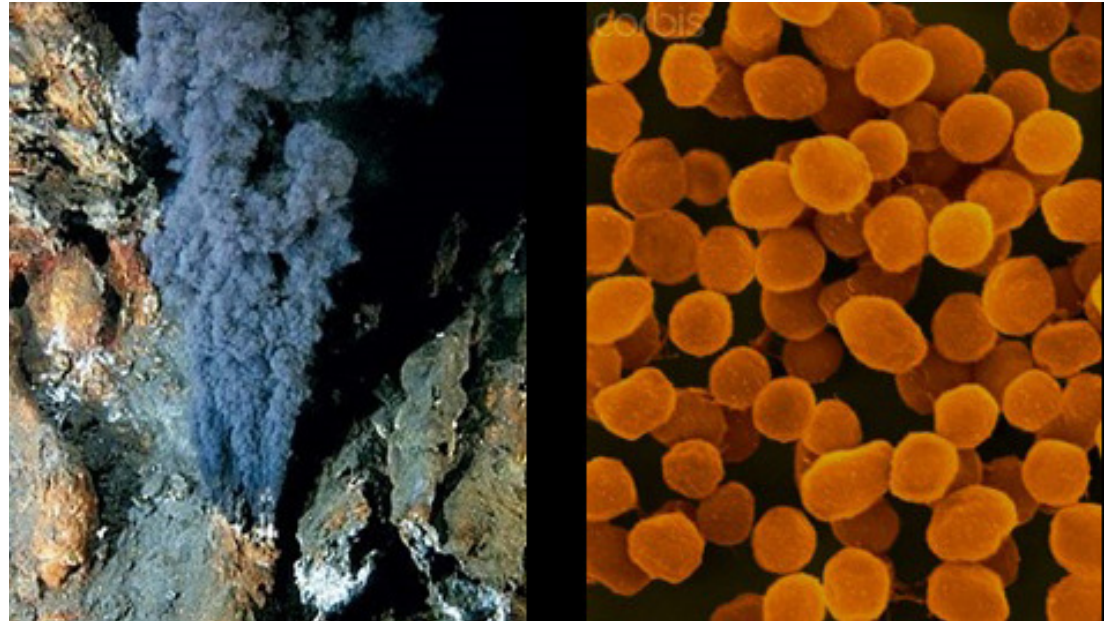




## Closest living relatives of LUCA



*clostridia*  
anaerobic bacteria  
(botulin, gangrene, tetanus)



Deep ocean vent - home to the extremophilic archeon  
*Methanococcus jannaschii*

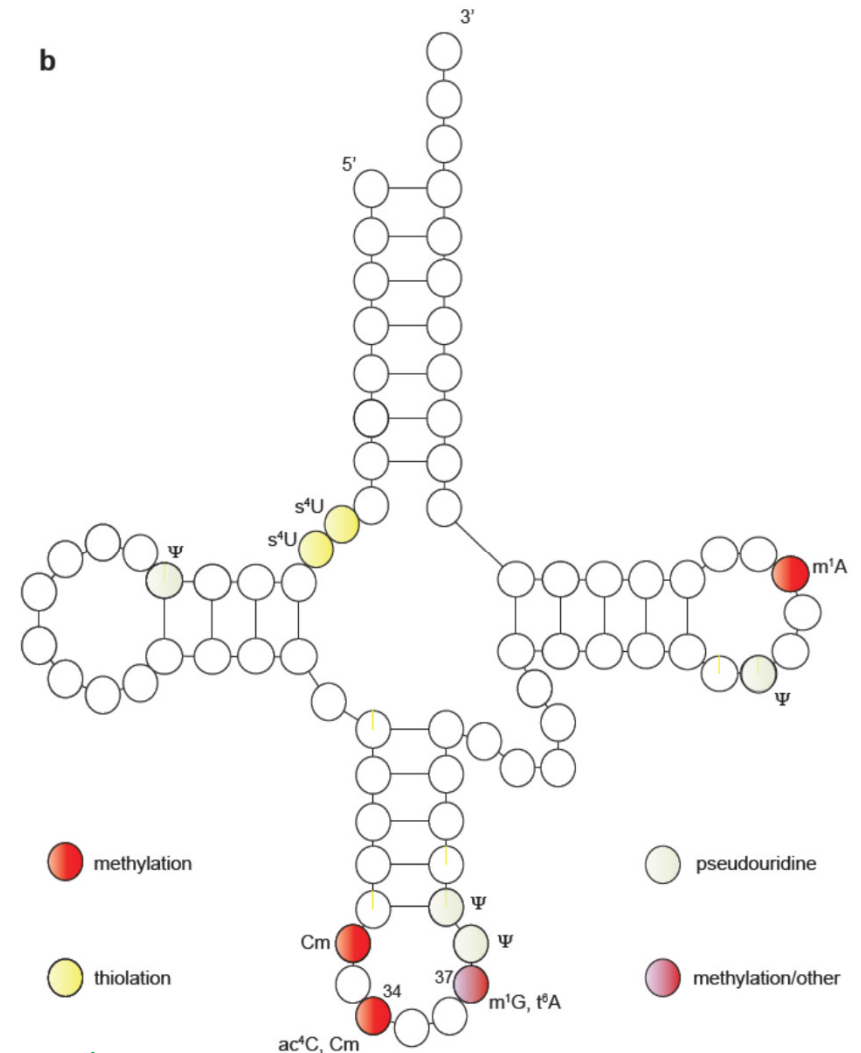
They use the WL pathway, abundant also today, some species can live from methyl groups (methane gas on marshes and wetlands), and they depend on H<sub>2</sub> (from geology or H<sub>2</sub>-producing fermentation)

Geological source of hydrogen: serpentinization (iron + hot water, anoxic)  $\text{Fe}^{2+} + \text{H}_2\text{O} \rightarrow \text{Fe}_3\text{O}_4 + \text{H}_2$

## Modified nucleosides and the genetic code

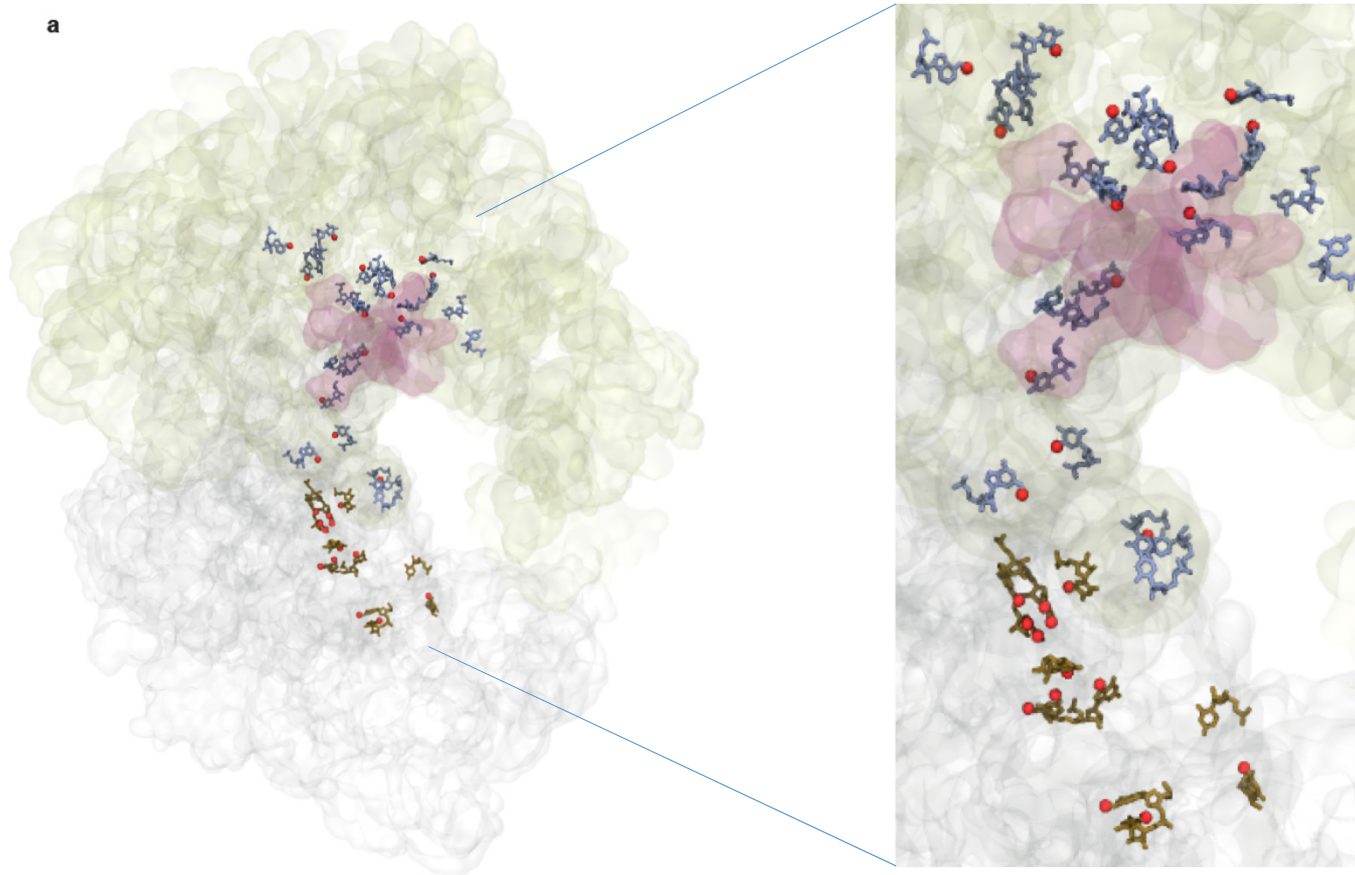
LUCA had also genes involved in RNA nucleoside modifications (mainly methylations and thiomethylations) still required today e.g. for the anticodon recognition process.

*Cloverleaf secondary structure representation of tRNA showing only those posttranscriptional nucleoside modifications that are conserved among bacteria and archaea in both identity and position. (5-methoxyuridine at position 34 in archaea has been disputed).*



M.C. Weiss et al. *Nature Microbiology*, 2016, Article 16116

## Modified nucleosides and the genetic code



M.C. Weiss et al. *Nature Microbiology*,  
2016, Article 16116

Structure of the *E. coli* ribosome (PDB ID: 4YBB), with the large and small subunits shown in green and silver, respectively. The peptidyl-transferase site is shaded pink. The modified nucleosides of 23S rRNA are depicted in icy blue, while in 16S rRNA they are ochre. Modification of C2501 to 5-hydroxycytidine is not present in the structure. Methyl group carbons are shown as red balls.



## ***Transition from the RNA world to LUCA***

***Ribozymes – self-acting → metabolic***

***Evolution of ribosome***

***Incorporation of aminoacids and peptides***

***The genetic code and archival storage***

***Enzyme-driven metabolism and membranes***

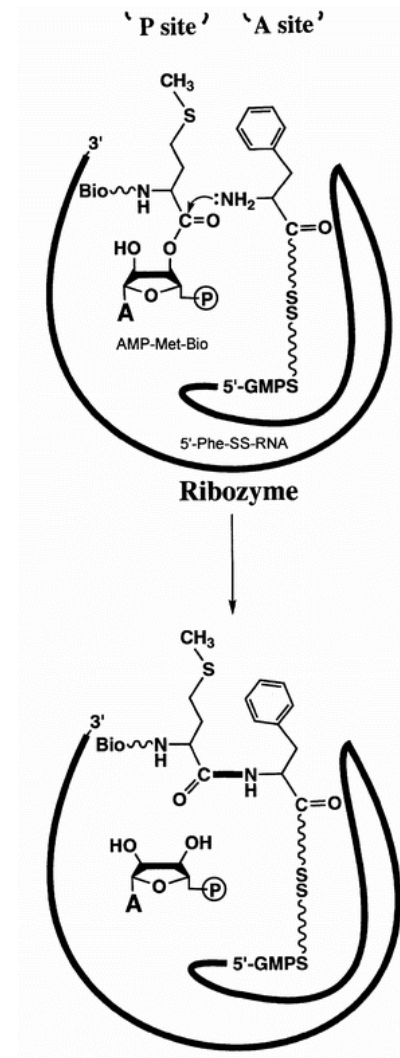
# Ribozymes

Initially only self-processing ribozymes (introns, RNAses) discovered.

1992 – first ribozyme isolated capable to cleave the bond of methionine with its tRNA (also the reverse reaction – transacylation – is catalysed)

1995 (Yarus) – a random RNA sequence found capable of attaching an activated aminoacid to itself

1997 (Szostak) – an RNA sequence that transfers one aminoacid to another one, forming a dipeptide → analogue of the peptidetransferase center of the ribosome



# Ribozymes

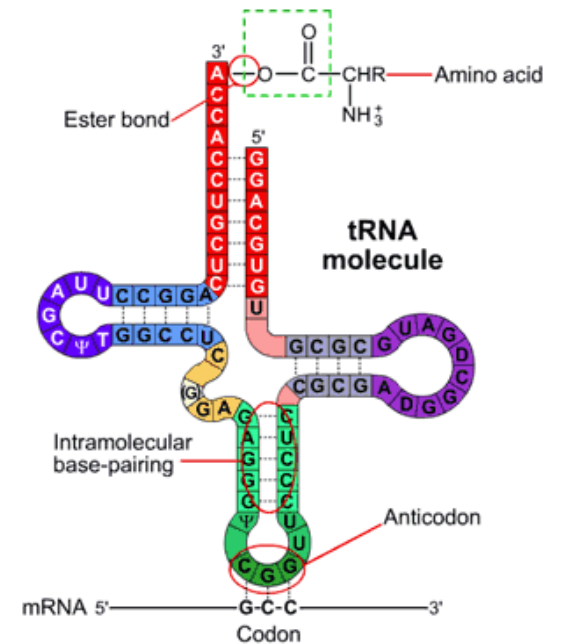
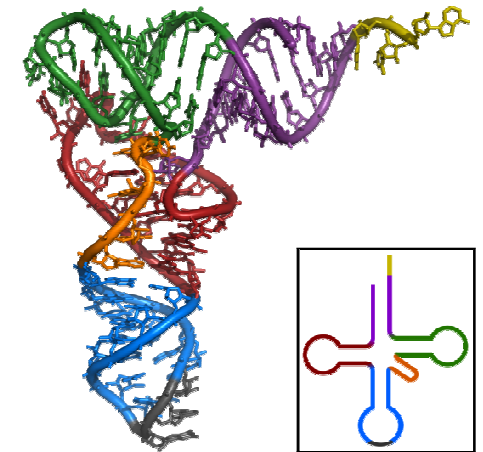
Ribozymes incorporate aminoacids to enhance their catalytic abilities

It opens ways to improved metabolism and provides evolutionary advantage in receiving energy from outside

Initially incorporation of aminoacids may have improved synthesis of nucleotides to produce more RNA

Primordial tRNAs were most likely self-charging, today special enzymes do it (tRNA synthethases)

Peptide chains increase in size, the RNA part decrease → non-covalent binding of nucleoside cofactors to contemporary enzymes

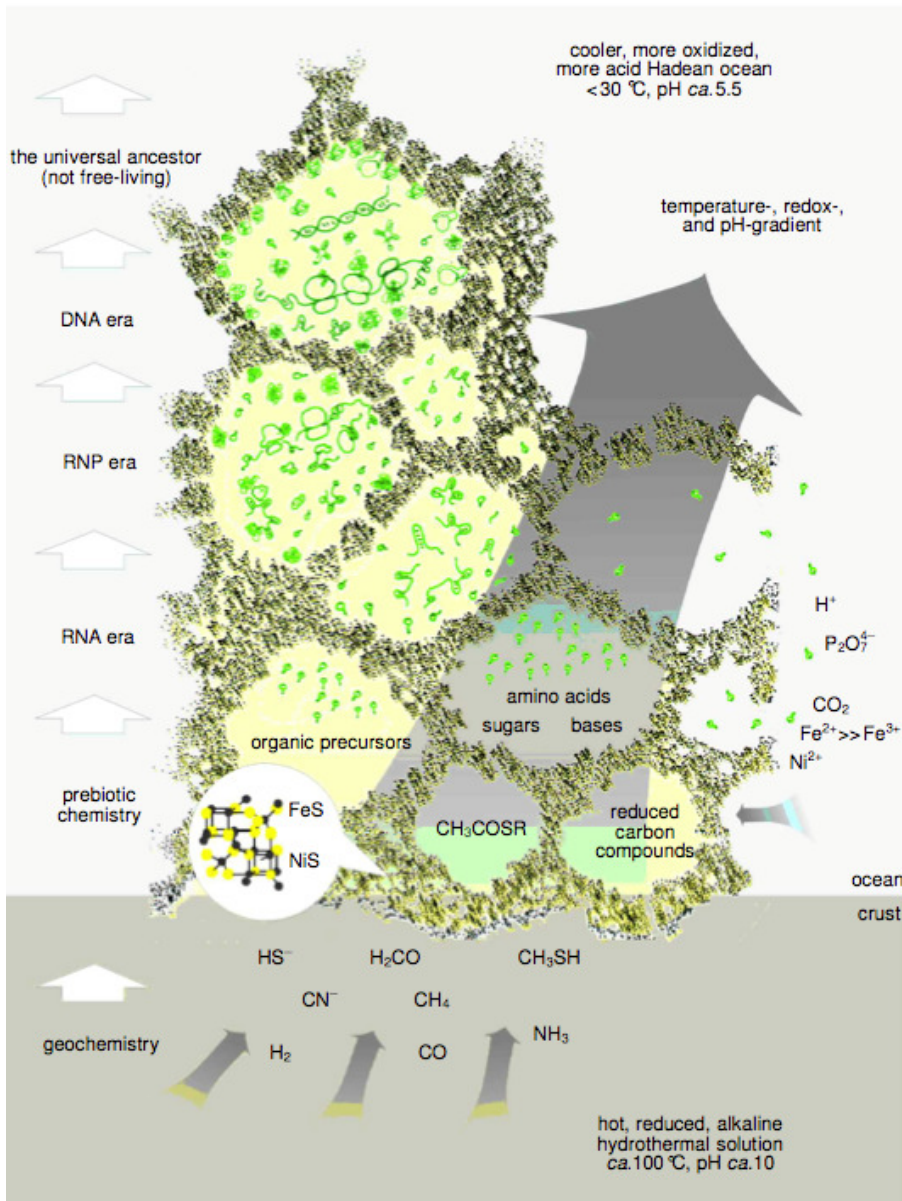


*aminoacyl-tRNA*

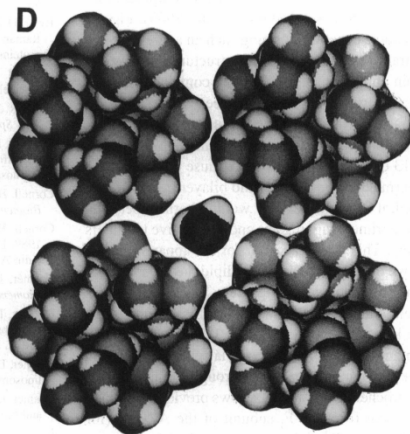
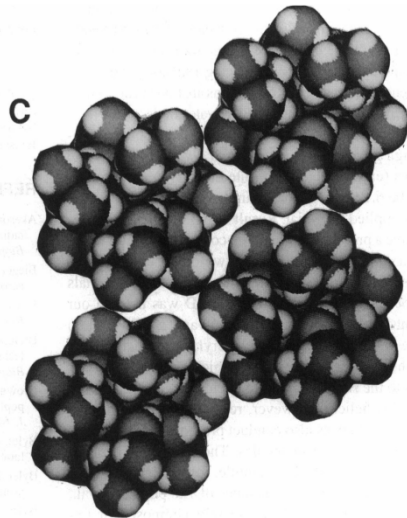
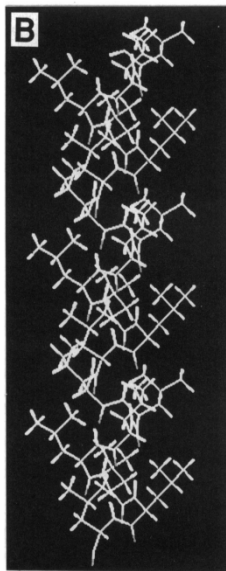
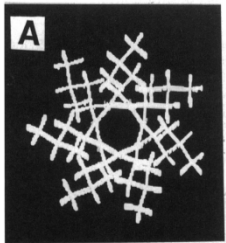
## Increasing metabolic complexity

Complex metabolic machinery closed in the same compartment that genetic polymers (RNA) which generated it.

*We don't see ribozyme-based metabolism today anymore, because protein catalysts (enzymes) for the same reactions are orders of magnitude faster than the ribozymes*



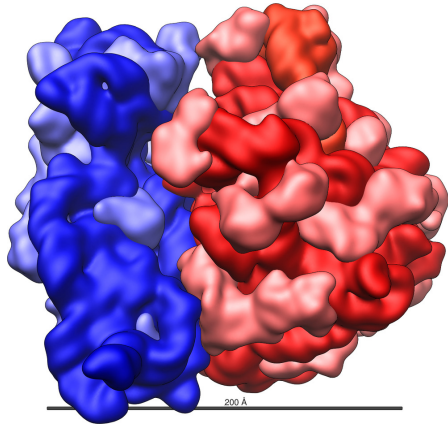
## Evolutionary advantage of proteins



Poly-alanine and poly-leucine form ion channels that selectively transport protons across lipid bilayers (not  $\text{Na}^+$  or  $\text{K}^+$ )

Short peptides with polar positively charged end (arginines) and unpolar Leu/Phe/Trp drive RNAs to membranes (Szostak)

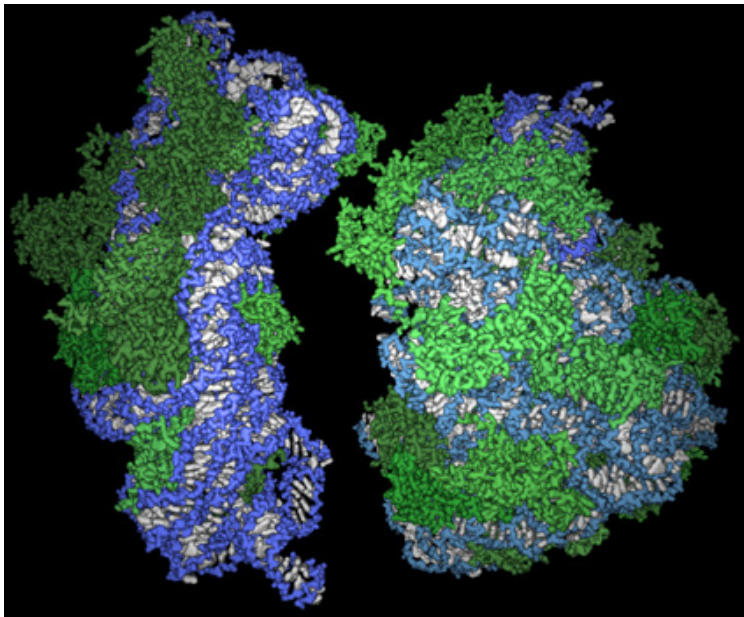
# Ribosome



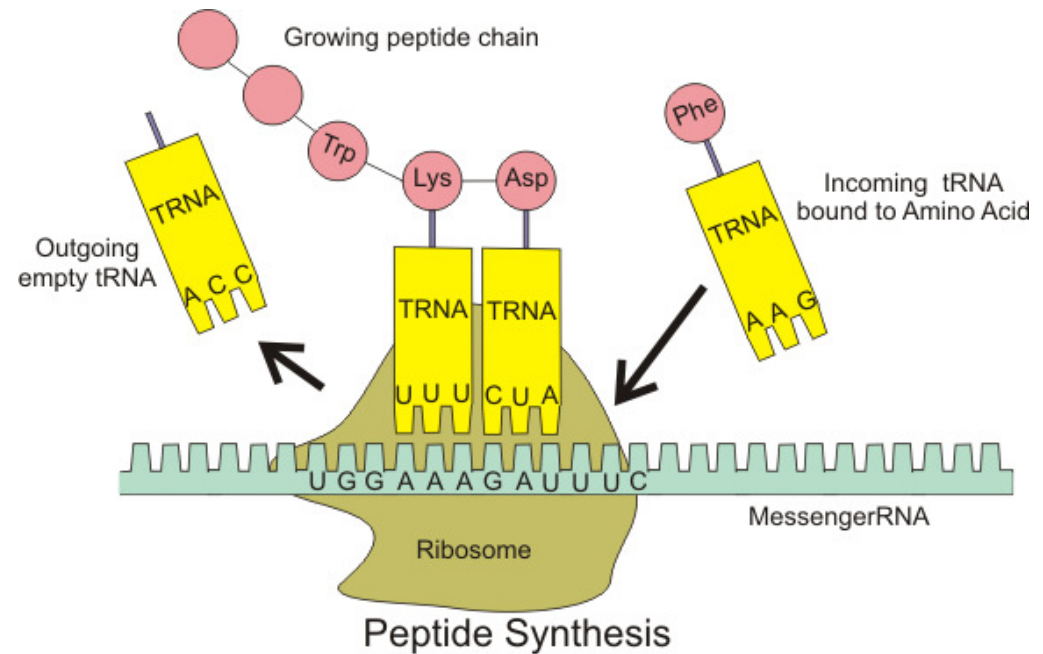
Every protein component of the ribosome can be removed without losing the activity.

Sequence-specific synthesis of proteins was invented late

Initially large subunit catalysed transacylations, later the small subunit used another RNA strand to ,guide' the new peptide growth in a sequence-specific manner (by codon-anticodon recognition). This strand (proto-mRNA) allowed tighter binding.

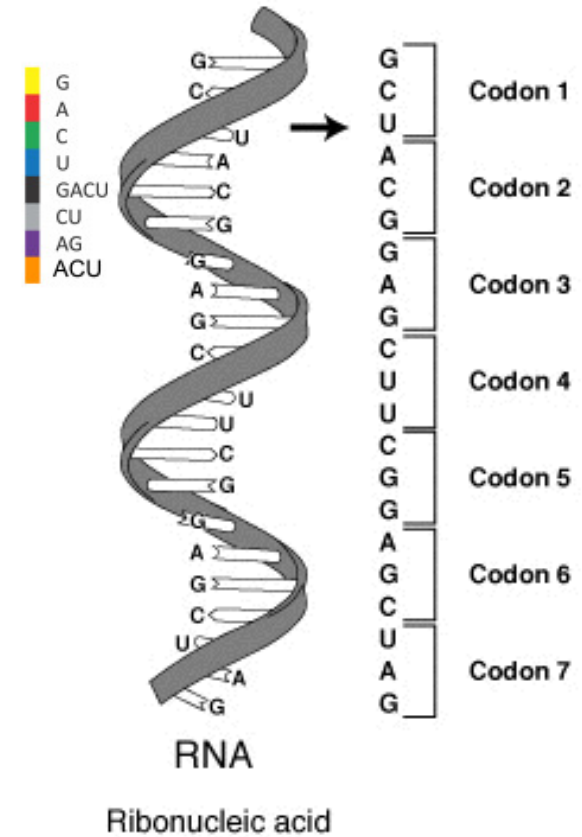
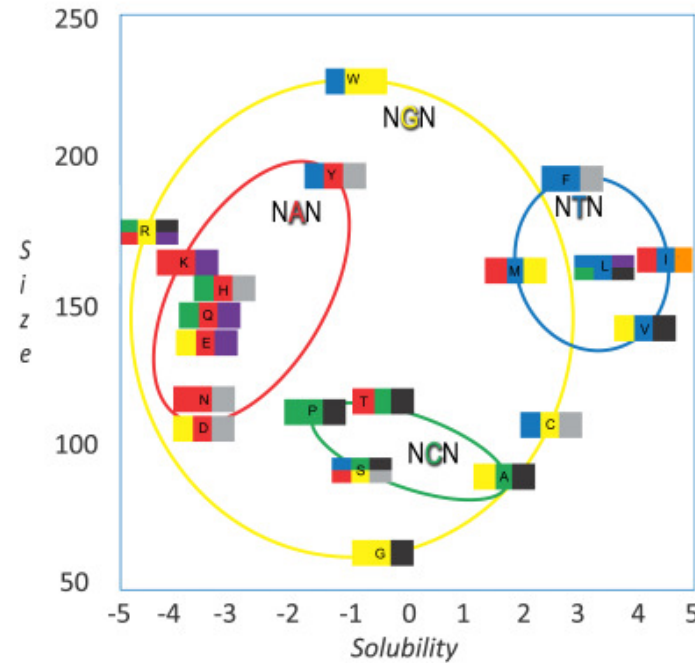


Ribosome: green - proteins, blue and white - RNA



# The genetic code

		Second letter				
		U	C	A	G	
First letter	U	UUU } Phe UUC } UUA } Leu UUG }	UCU } Ser UCC } UCA } UCG }	UAU } Tyr UAC } UAA Stop UAG Stop	UGU } Cys UGC } UGA Stop UGG Trp	U C A G
	C	CUU } Leu CUC } CUA } CUG }	CCU } Pro CCC } CCA } CCG }	CAU } His CAC } CAA } Gln CAG }	CGU } Arg CGC } CGA } CGG }	U C A G
	A	AUU } Ile AUC } AUA } AUG Met	ACU } Thr ACC } ACA } ACG }	AAU } Asn AAC } AAA } Lys AAG }	AGU } Ser AGC } AGA } Arg AGG }	U C A G
	G	GUU } Val GUC } GUA } GUG }	GCU } Ala GCC } GCA } GCG }	GAU } Asp GAC } GAA } Glu GAG }	GGU } Gly GGC } GGA } GGG }	U C A G



# The genetic code

## Examples of notable Mutations

ΔF508 deletion in cystic fibrosis

		2nd base			
		U	C	A	G
1st base	U	UUU (Phe/F) Phenylalanine	UCU (Ser/S) Serine	UAU (Tyr/Y) Tyrosine	UGU (Cys/C) Cysteine
		UUC (Phe/F) Phenylalanine	UCC (Ser/S) Serine	UAC (Tyr/Y) Tyrosine	UGC (Cys/C) Cysteine
		UUA (Leu/L) Leucine	UCA (Ser/S) Serine	UAA Ochre (Stop)	UGA Opal (Stop)
		UUG (Leu/L) Leucine	UCG (Ser/S) Serine	UAG Amber (Stop)	UGG (Trp/W) Tryptophan
	C	CUU (Leu/L) Leucine	CCU (Pro/P) Proline	CAU (His/H) Histidine	CGU (Arg/R) Arginine
		CUC (Leu/L) Leucine	CCC (Pro/P) Proline	CAC (His/H) Histidine	CGC (Arg/R) Arginine
		CUA (Leu/L) Leucine	CCA (Pro/P) Proline	CAA (Gln/Q) Glutamine	CGA (Arg/R) Arginine
		CUG (Leu/L) Leucine	CCG (Pro/P) Proline	CAG (Gln/Q) Glutamine	CGG (Arg/R) Arginine
	A	AUU (Ile/I) Isoleucine	ACU (Thr/T) Threonine	AAU (Asn/N) Asparagine	AGU (Ser/S) Serine
		AUC (Ile/I) Isoleucine	ACC (Thr/T) Threonine	AAC (Asn/N) Asparagine	AGC (Ser/S) Serine
		AUA (Ile/I) Isoleucine	ACA (Thr/T) Threonine	AAA (Lys/K) Lysine	AGA (Arg/R) Arginine
		AUG (Met/M) Methionine	ACG (Thr/T) Threonine	AAG (Lys/K) Lysine	AGG (Arg/R) Arginine
G	GUU (Val/V) Valine	GCU (Ala/A) Alanine	GAU (Asp/D) Aspartic acid	GGU (Gly/G) Glycine	
	GUC (Val/V) Valine	GCC (Ala/A) Alanine	GAC (Asp/D) Aspartic acid	GGC (Gly/G) Glycine	
	GUA (Val/V) Valine	GCA (Ala/A) Alanine	GAA (Glu/E) Glutamic acid	GGA (Gly/G) Glycine	
	GUG (Val/V) Valine	GCG (Ala/A) Alanine	GAG (Glu/E) Glutamic acid	GGG (Gly/G) Glycine	

Selection of notable mutations, ordered in a standard table of the genetic code of amino acids.

Clinically important missense mutations generally change the properties of the coded amino acid residue between being basic, acidic, polar or nonpolar, while nonsense mutations result in a stop codon.

**Amino acids**

- Basic
- Acidic
- Polar
- Nonpolar (hydrophobic)

Fragile X Syndrome

**Polyglutamine (PolyQ) Diseases**

- Huntington's disease
- Spinocerebellar ataxia (SCA) (most types)
- Spinobulbar muscular atrophy (Kennedy disease)
- Dentatorubral-pallidolusian atrophy

**Mutation type**

- Trinucleotide repeat
- Deletion
- Missense
- Nonsense

3rd base in each row

- Myotonic dystrophy  
- SCA 8

Prostate cancer

Colorectal cancer

Sickle-cell disease

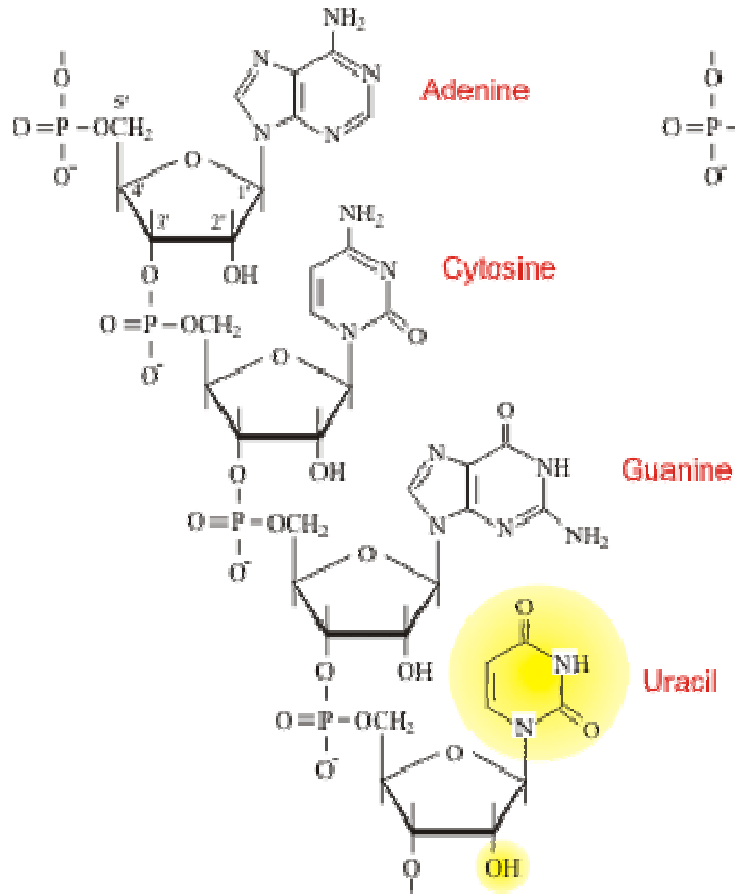
Friedreich's ataxia

β-Thalassemia

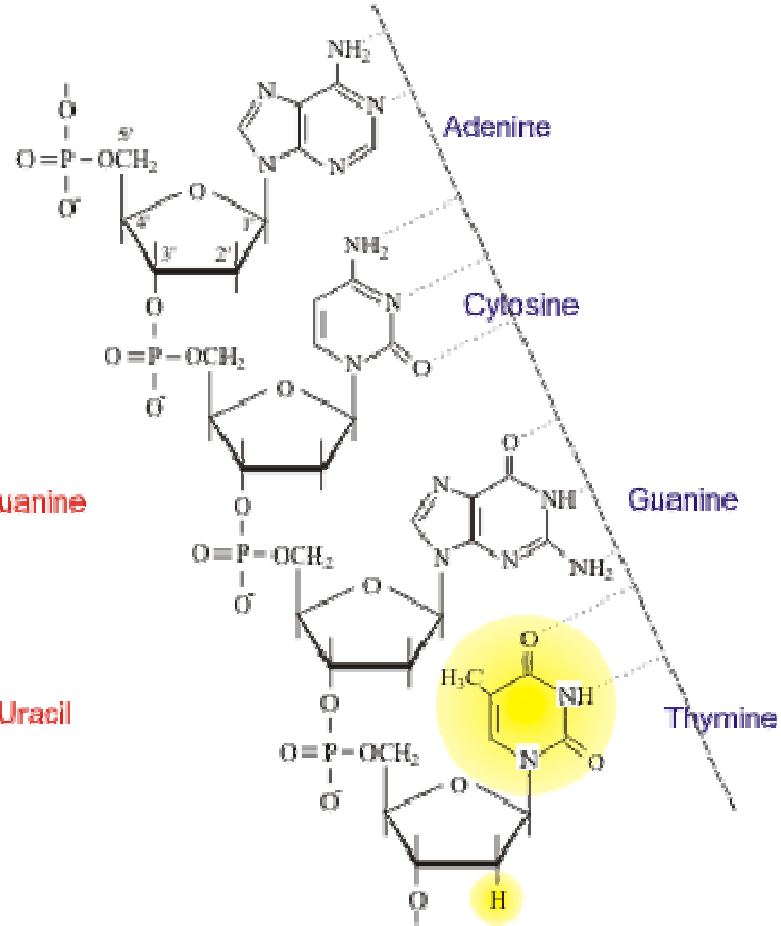
McArdle's disease



# The origin of DNA

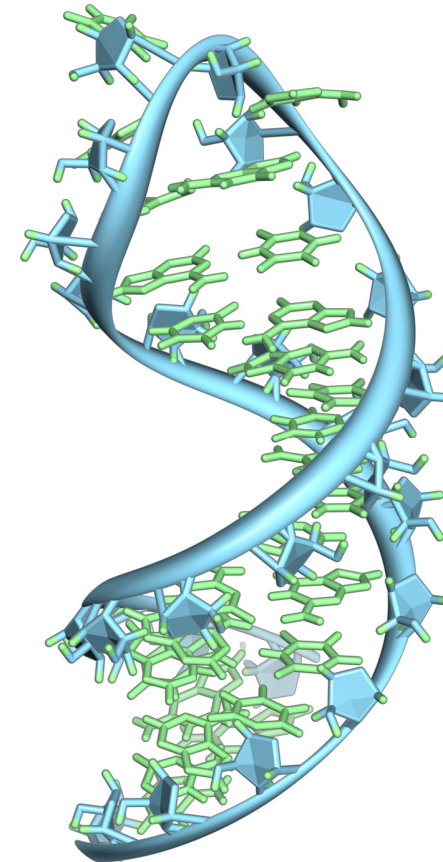
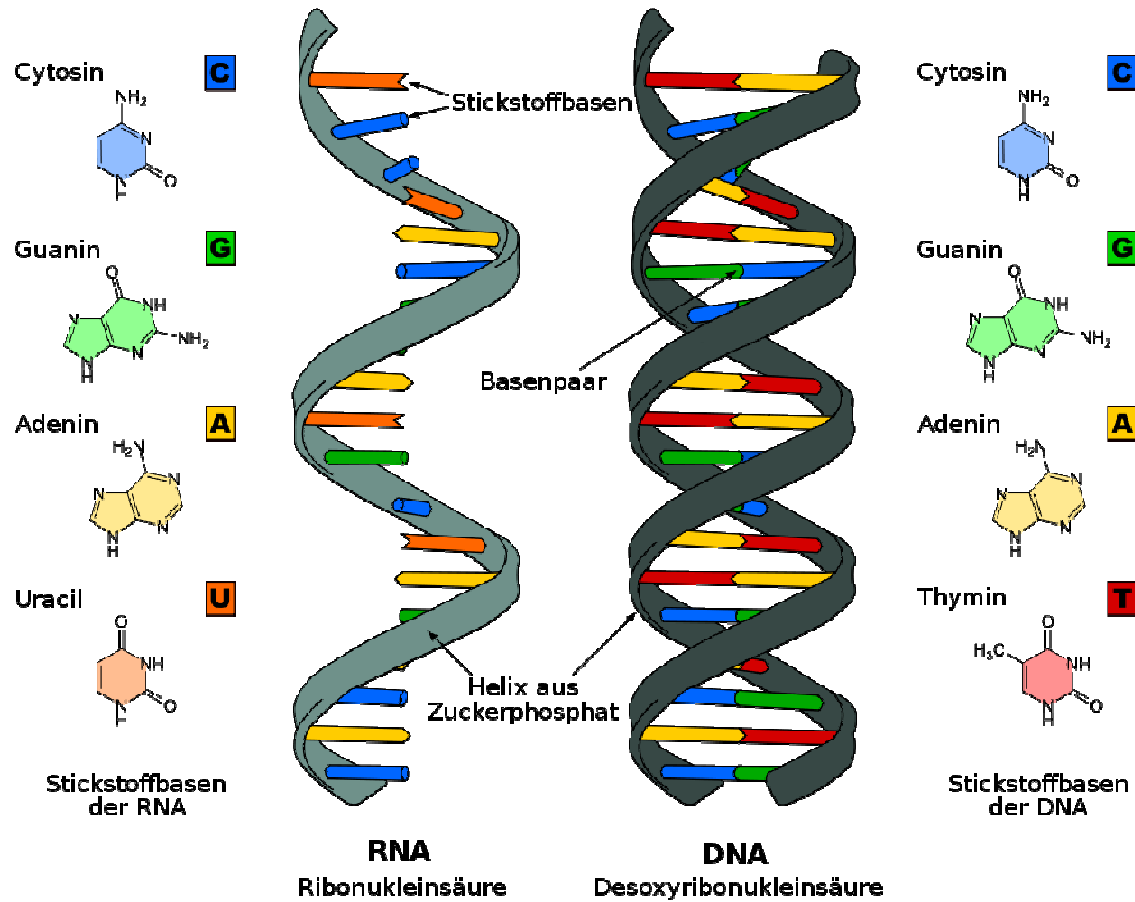


RNA

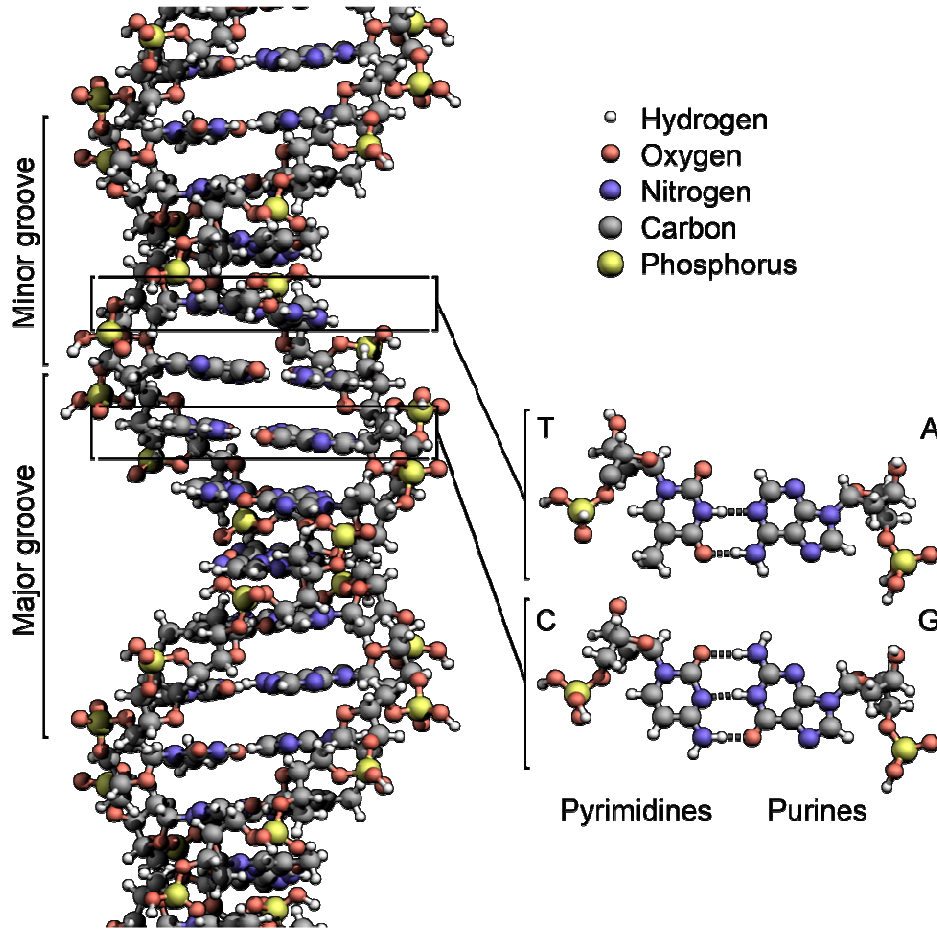


DNA

# The origin of DNA



# The origin of DNA



Maximal size of RNA-based genome: 3000-5000 bases  
(HIV, West Nile virus)

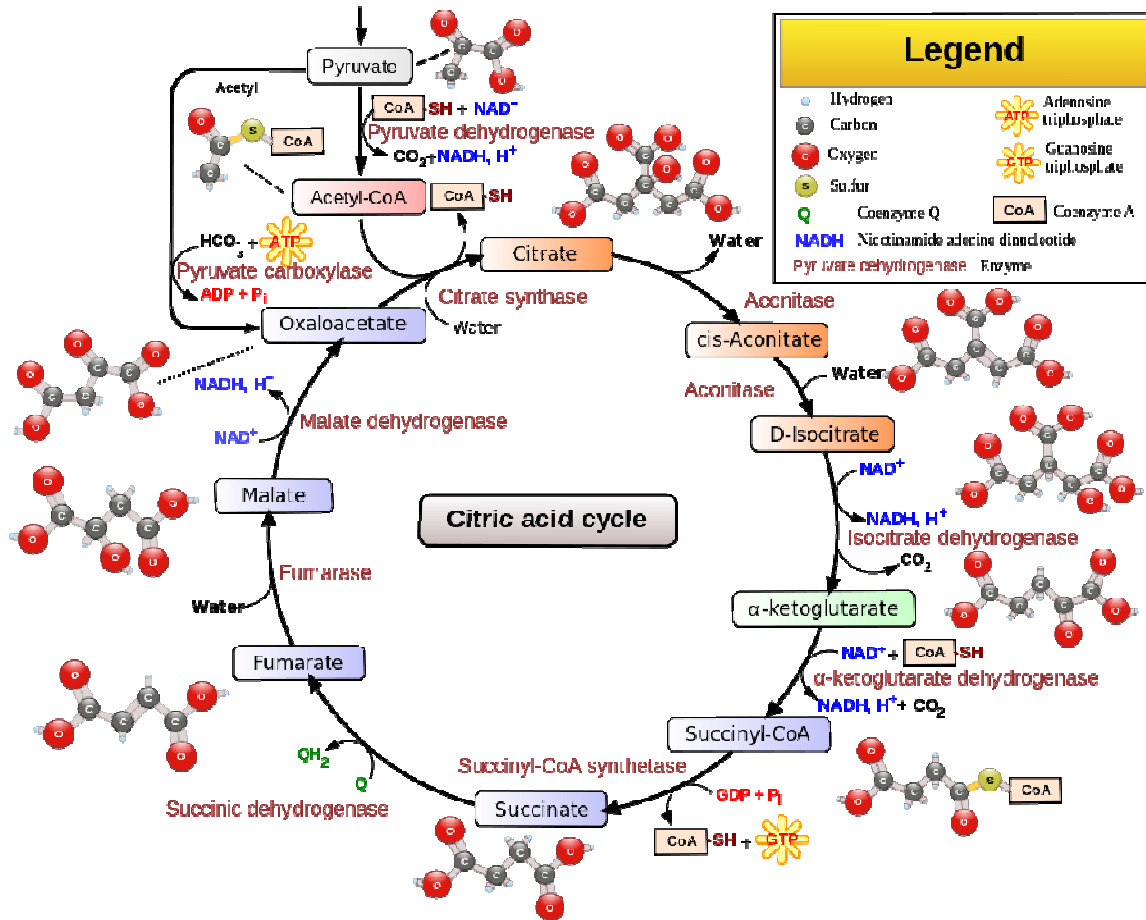
Reason: above that, statistically certain to generate at least one self-cleaving RNA sequence (ribozyme)

Maximal DNA size – unlimited

- no self-cleaving DNAzymes,
- tight storage as dsDNA,
- methylated uracil (thymine) → no accidental C-to-U mutations

# Metabolic pathways

## Citric acid cycle



Oxidative, not present in Archaea, most likely absent in LUCA

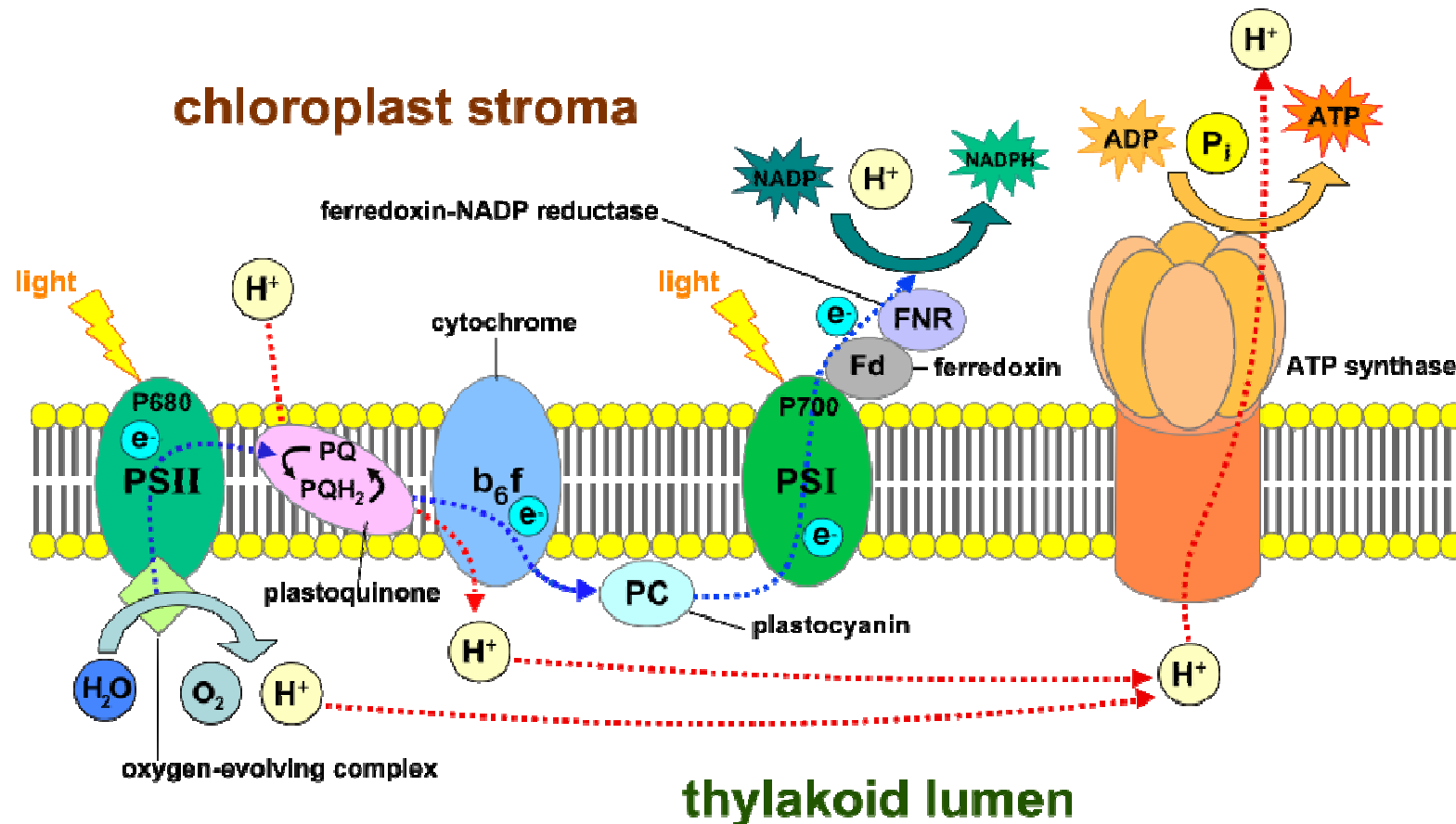
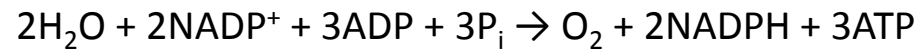
## Light harvesting - photosynthesis



PS I from green sulfur bacteria *Chlorobiaceae*

## Light harvesting - photosynthesis

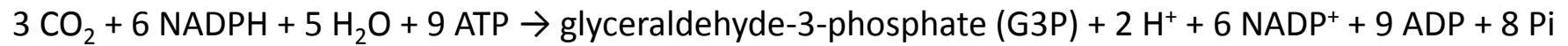
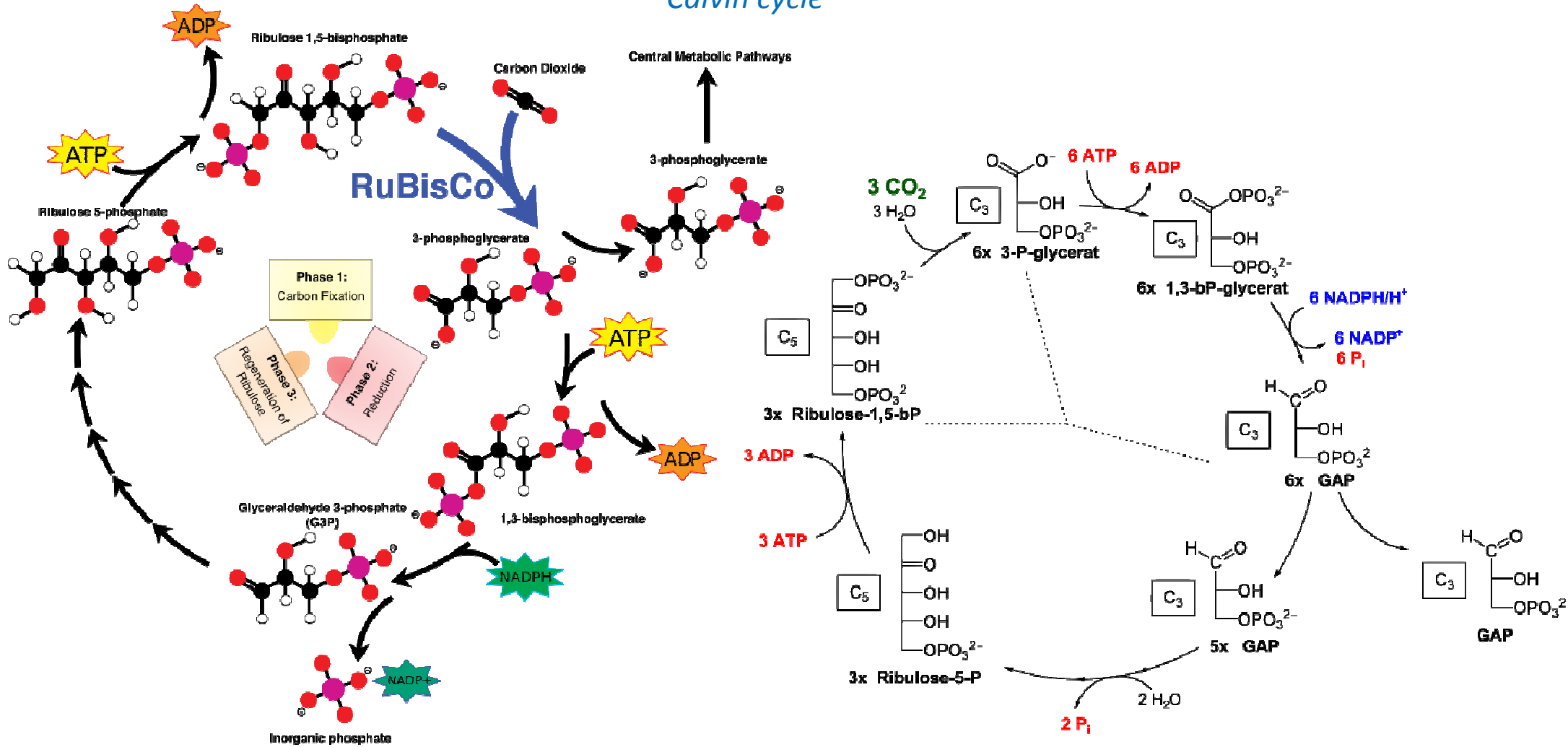
Light-dependent reactions of photosynthesis at the thylakoid membrane



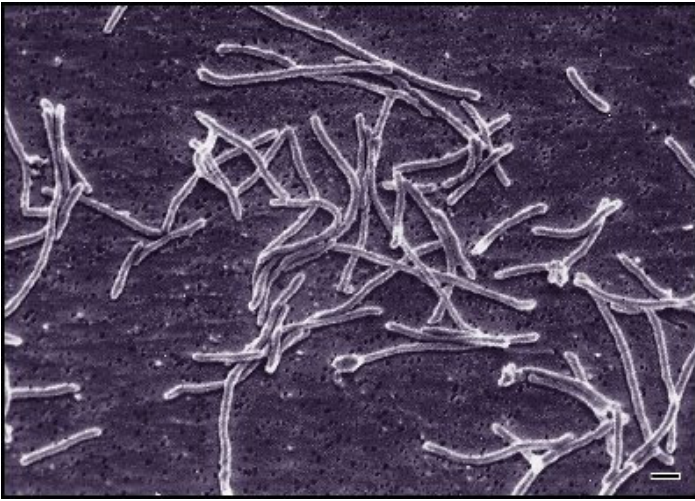
Water difficult to oxidize. Only combination of two photosystems provides enough electrochemical potential.

# Metabolic pathways

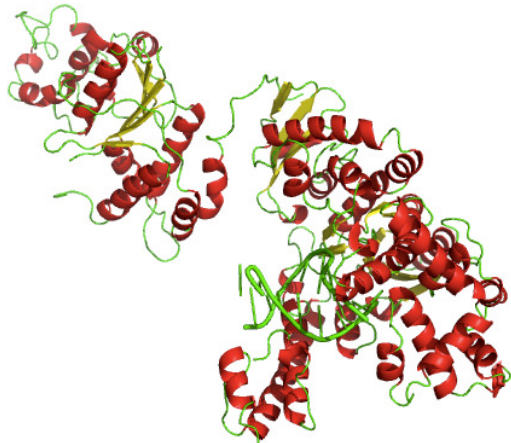
## Calvin cycle



## *Thermophiles*



*Thermus aquaticus*



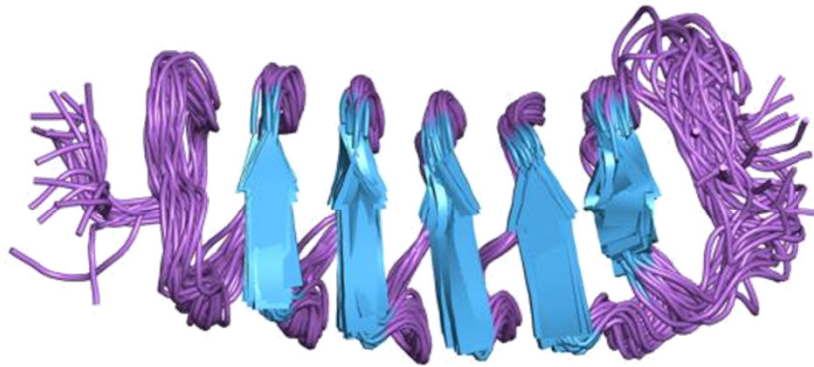
3D structure of Taq Polymerase.



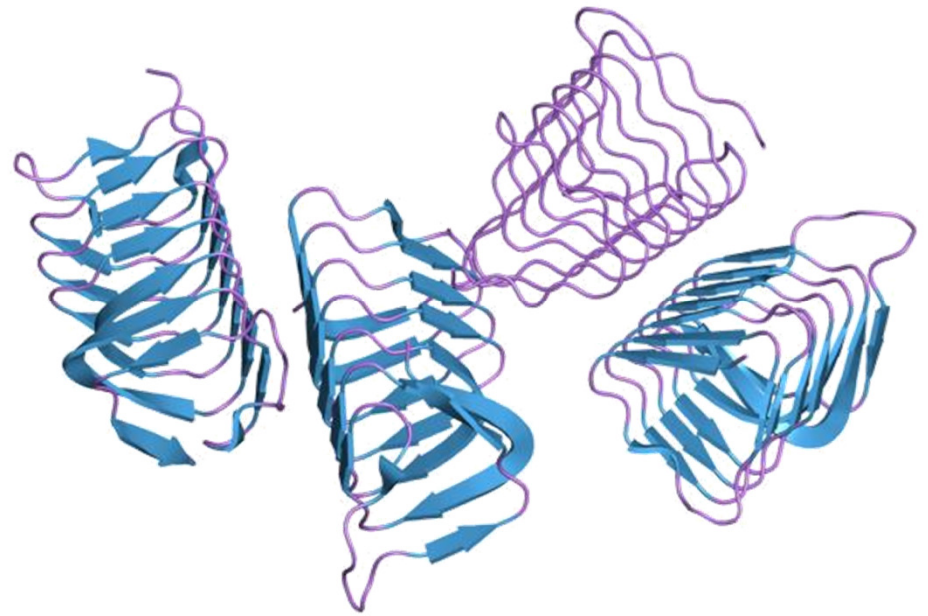
Hot springs with algae and bacteria in Yellowstone National Park



## Cold adaptation

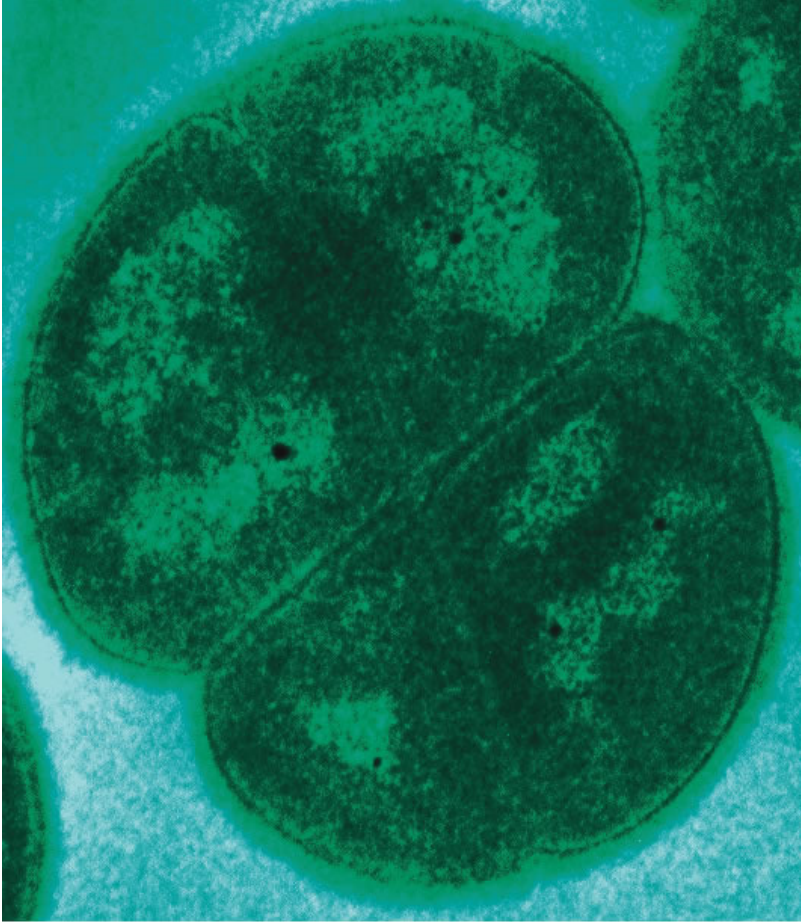


Structure of the *Tenebrio molitor* beta-helical antifreeze protein



Structure of *Choristoneura fumiferana* (spruce budworm) beta-helical antifreeze protein

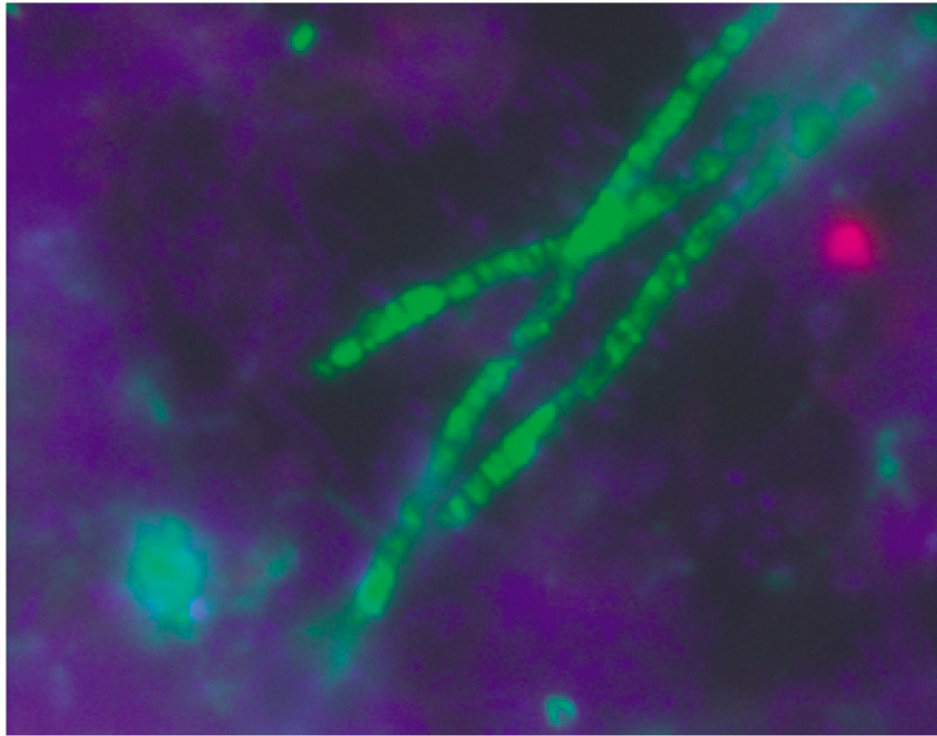
## *Drought, salinity, radiation*



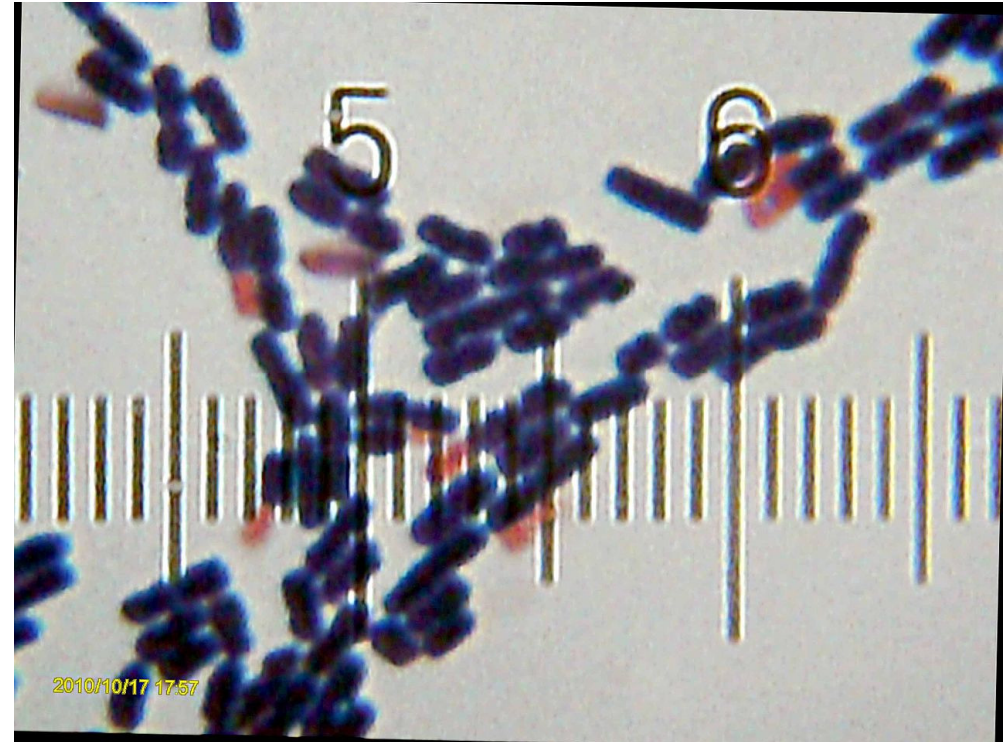
A tetrad of *D. radiodurans*

Efficient DNA damage repair,  
Trehalose as the main sugar – glass solid, no crystallization

## *Acid, base*



*Acidobacterium*



A typical *bacillus* culture. Many alkaliphiles possess a *bacillus* morphology



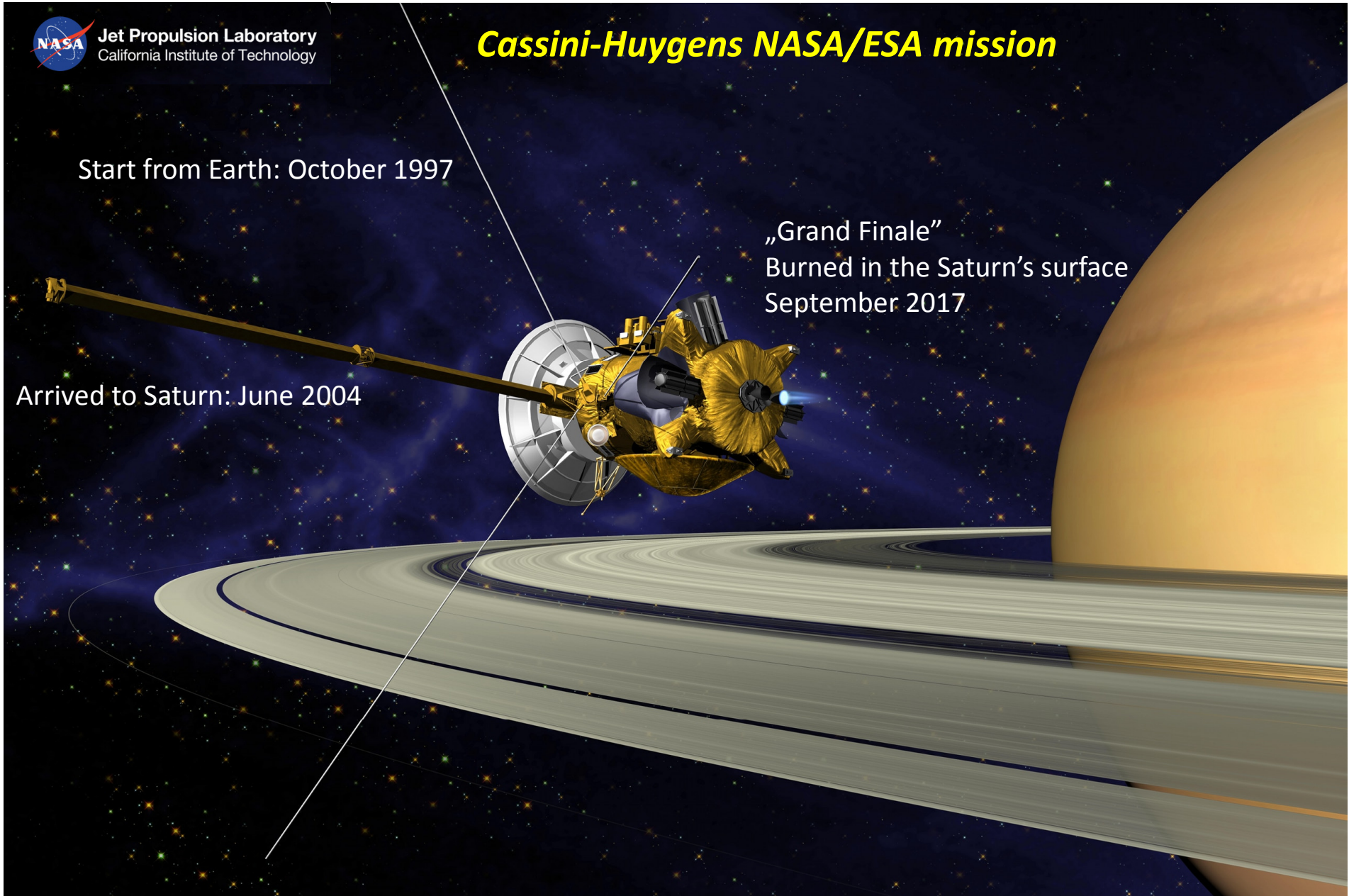
Jet Propulsion Laboratory  
California Institute of Technology

## *Cassini-Huygens NASA/ESA mission*

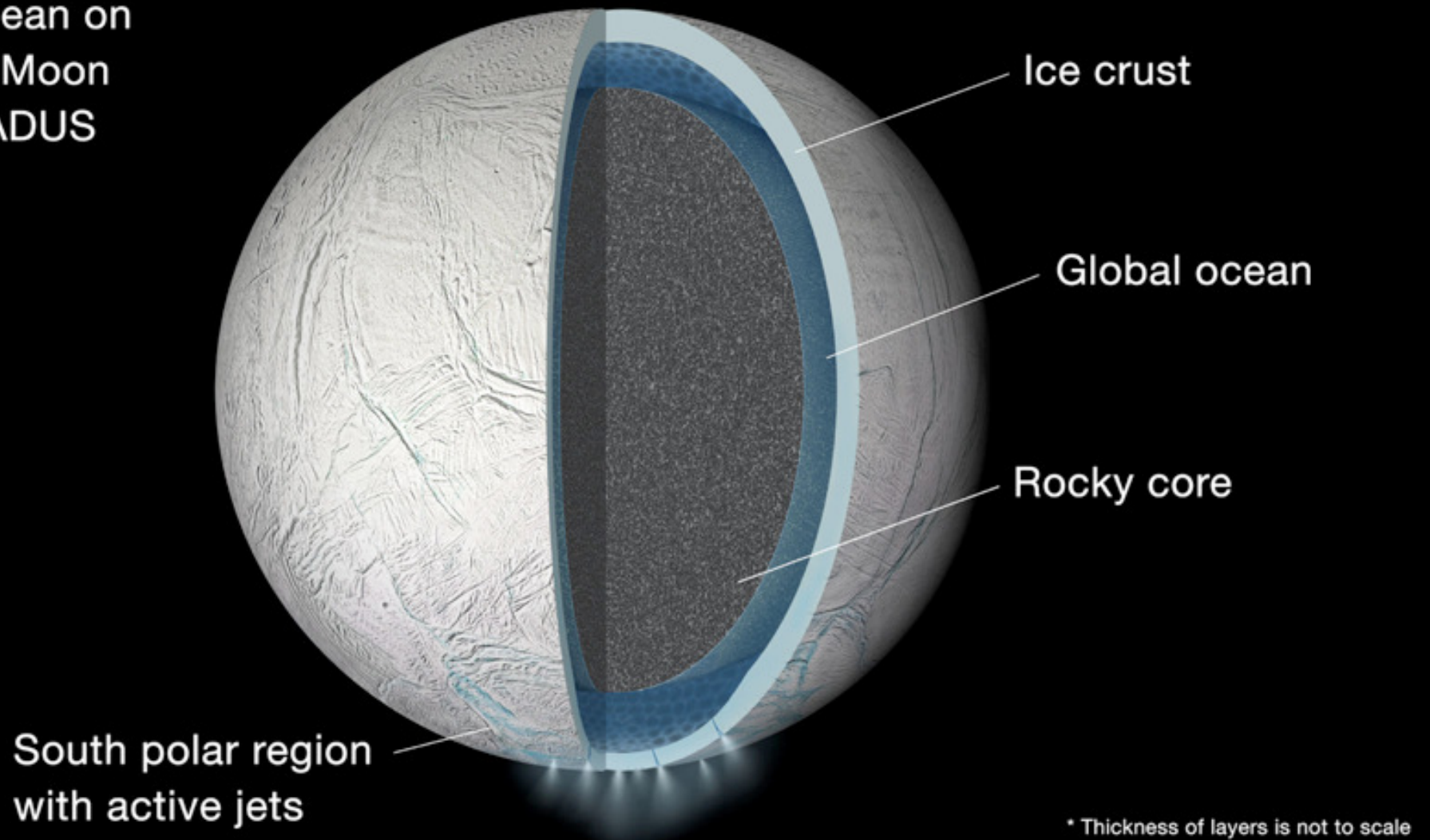
Start from Earth: October 1997

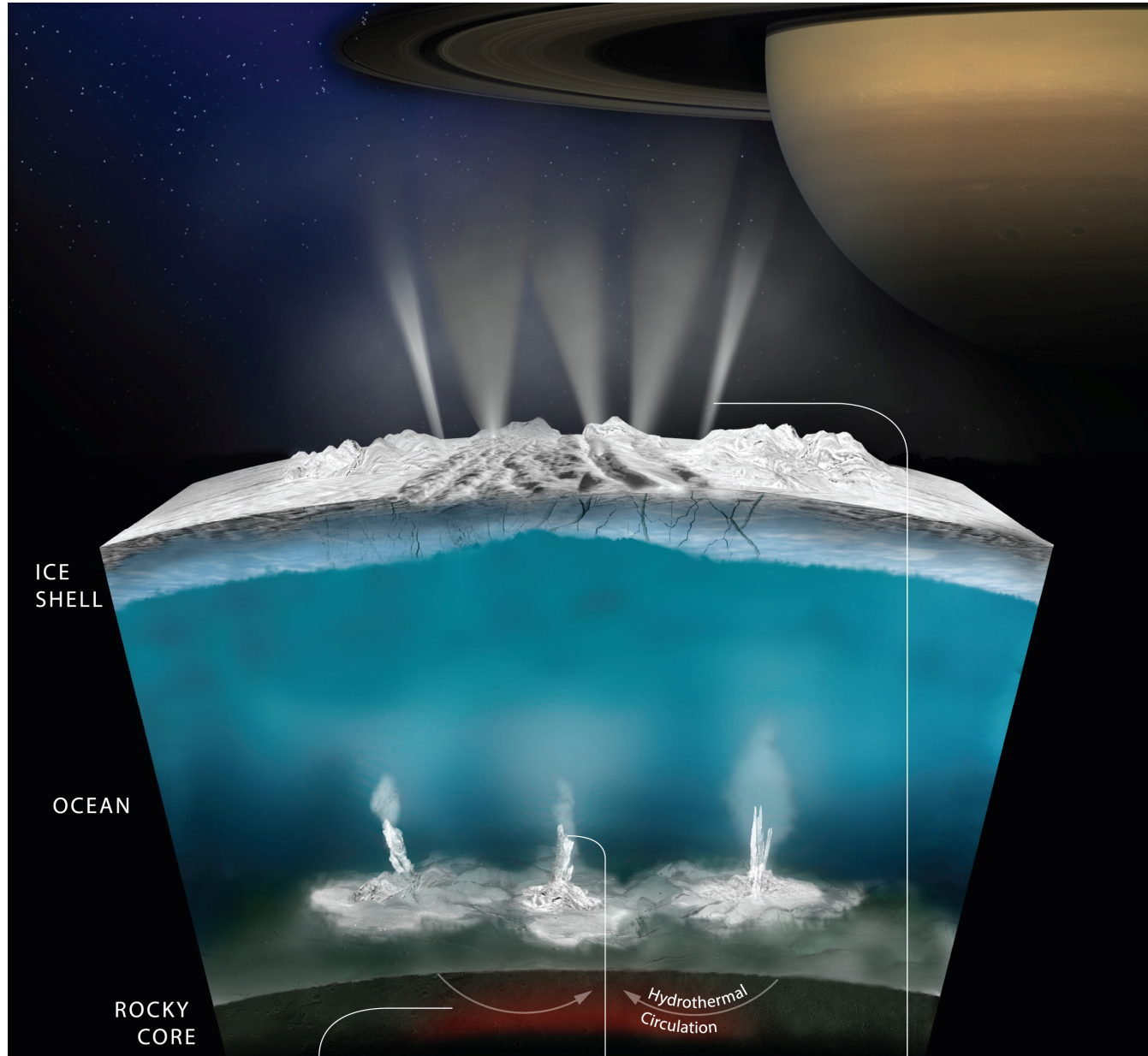
„Grand Finale“  
Burned in the Saturn's surface  
September 2017

Arrived to Saturn: June 2004



Global Ocean on  
Saturn's Moon  
ENCELADUS

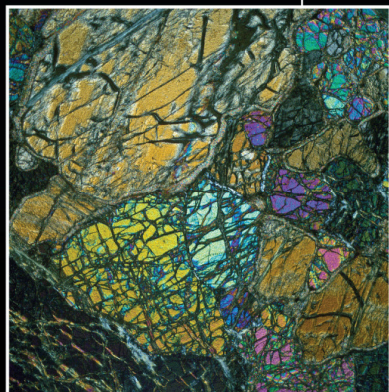




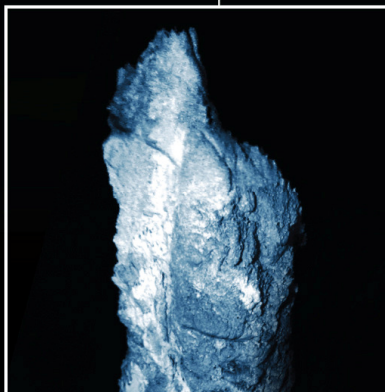
OCEAN

ROCKY  
CORE

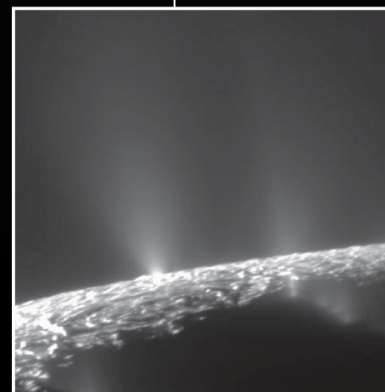
Hydrothermal  
Circulation



WATER-ROCK REACTIONS



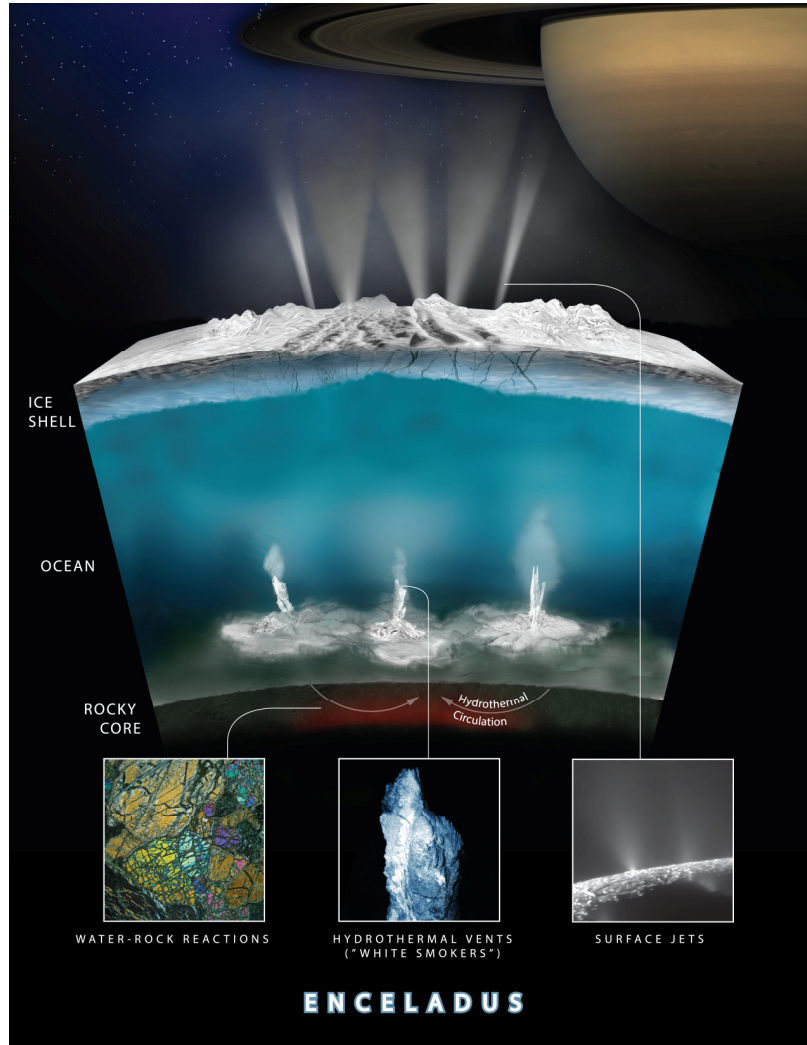
HYDROTHERMAL VENTS  
("WHITE SMOKERS")



SURFACE JETS

# ENCELADUS

# Can (and does?) Enceladus host (microbial) life?



## Fuels for life on one of Saturn's moons

NASA's Cassini spacecraft has discovered hydrogen and carbon dioxide erupting from Saturn's moon Enceladus – critical ingredients that sustain microbial life in extreme environments on Earth

**ENCELADUS**  
Diameter: 504km

**CASSINI**

2015: During Cassini's deepest-ever dive into plume of spray, instruments detected presence of organic chemicals in plume vapour

Geysers erupting from cracks in south polar region

**Methanogenesis:** Process by which microorganisms on Earth obtain energy by using hydrogen to produce methane from carbon dioxide

Hydrogen and carbon dioxide...  
...produce methane and water

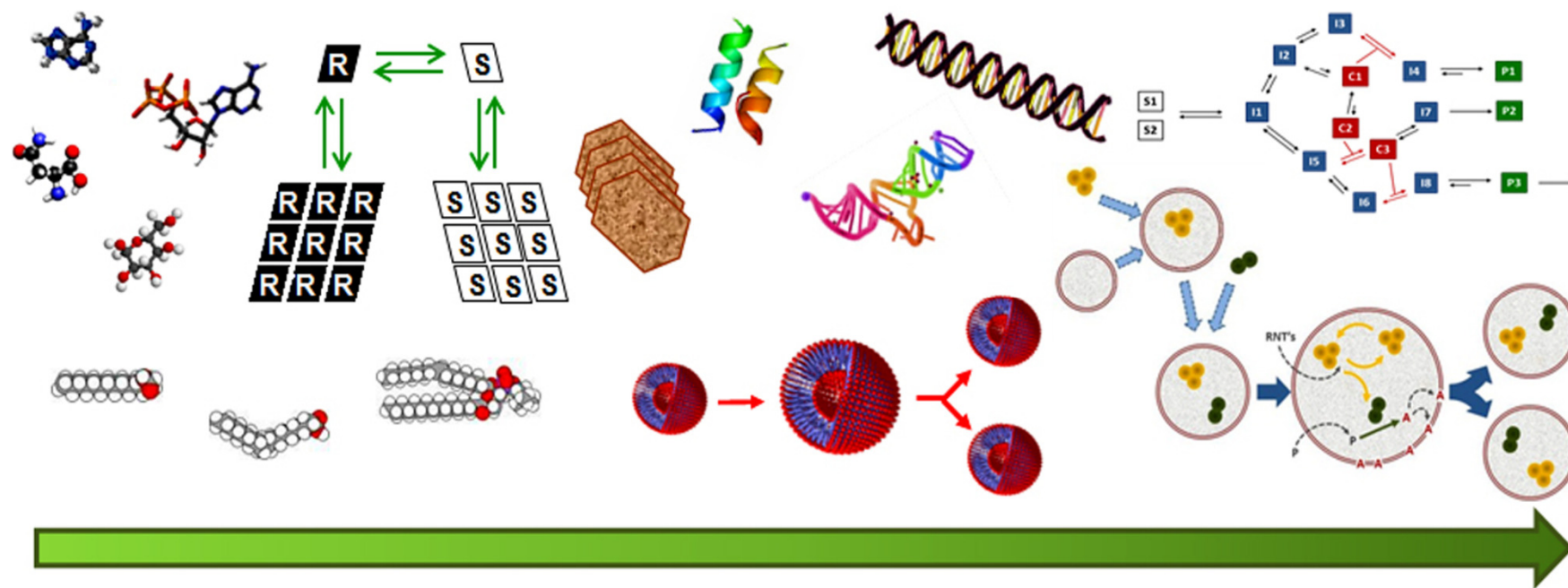
**Hydrothermal vent:** Molecular hydrogen most likely formed by chemical reactions between hot rocks and water in ocean

Sources: Science, Jet Propulsion Laboratory, Hunter Waite and colleagues © GRAPHIC NEWS



## *The molecular origins of life – important stages*

biomolecules – biopolymers – self-replication – metabolism - compartmentalization



Increasing complexity from molecules to systems