## The Transition to Life

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Interacting Chemical  $\longrightarrow$  Reproduction of Organisms

### Based on Simplest Life Now:

#### Need:

- 1. Nucleic Acids
- 2. Proteins
- 3. Lipids
- 4. Carbohydrates (Pigments)

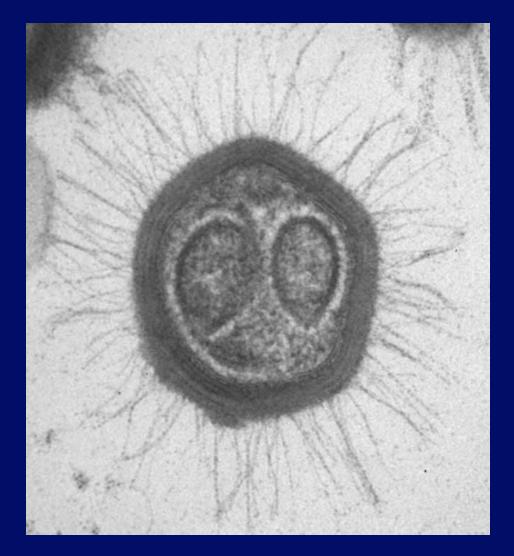
Replicable Information Enzymes (Catalysts) Membranes (Enclosure) Energy Storage (Energy Conversion)

Too much to ask of chemical evolution ⇒ Protolife?

### Mimiviruses: A Model for protolife?

- A very large virus (mimivirus) was discovered in 2003
- Now part of a group of such viruses
- Both RNA and DNA
- More DNA than some bacteria (> 1000 genes)
- Genes for translation, DNA repair enzymes
- Still needs ribosome of a host cell
- Leading to reevaluation of viruses
- May be ancient lineages
  - Precursors to bacteria, or eukaryotes
  - Controversial

# Image of Mimivirus



### Protolife

1. "Virus" equivalent in complexity, but free-living

Protein + Nucleic Acid + Supply by Environment

- 2. Protein Protolife Protein → Self Replication?
- 3. Nucleic Acid Protolife
   RNA → Self Catalysis?

4. Something Else
Minerals
Clay Layers
Mineral - Molecule
Pyrite
Thioesters

**Genetic Takeover** 

 $? \longrightarrow \mathsf{RNA} \longrightarrow \mathsf{DNA}$ 

### **Protein-based Protolife**

- Experiments by Sidney Fox
- Amino acids + dry heat led to proteinoids
- Addition of water produced microspheres
- They can grow, bud, form chains, divide
- But no nucleic acids, so not reproduction
- Could be model for protocells
- Consider "evolution" to life later

### **Picture of Proteinoid Microspheres**

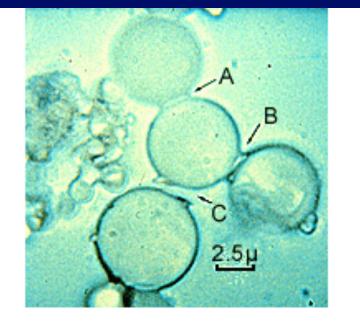


FIGURE 5.15 — Photograph of proteinoid microspheres produced by repeated energizing and dehydrating the primordial soup. The main features of this figure can be simulated by shaking a mixture of oil and water and watching the globs of oil cluster on the surface of the water. Seen here through a microscope, each microsphere contains a large\ concentration of amino acids. (The scale shown, 2.5 microns, equals 2.5x10-4 cm.) (Sidney Fox)

#### Nucleic Acid Based Protolife

RNA — Genes — Protein — A Cells Self-replicating RNA molecules Experiment by Sol Spiegelman RNA from Q<sub>β</sub> Virus - parasite on bacteria Injects RNA - Bacterium makes replicase **Enzyme to Replicate RNA** RNA multiplies, using activated nucleotides in bacterium to copy RNA and make new viruses

#### In Test Tube: Template RNA, Replicase, Activated Nucleotides (ATP, CTP, GTP, UTP)

⇒ RNA copied **without** machinery of cell

Variation: No template RNA

Replicase made RNA from nucleotides

Protein

Manfred Eigen - further experiments with RNA in test tube:

Mutant RNA strands compete Degrade to smallest (~ 200 nucleotides) RNA that replicase could recognize (Monster - Selfish RNA)

RNA can do self-catalysis in some cases Could this have led to self replication?

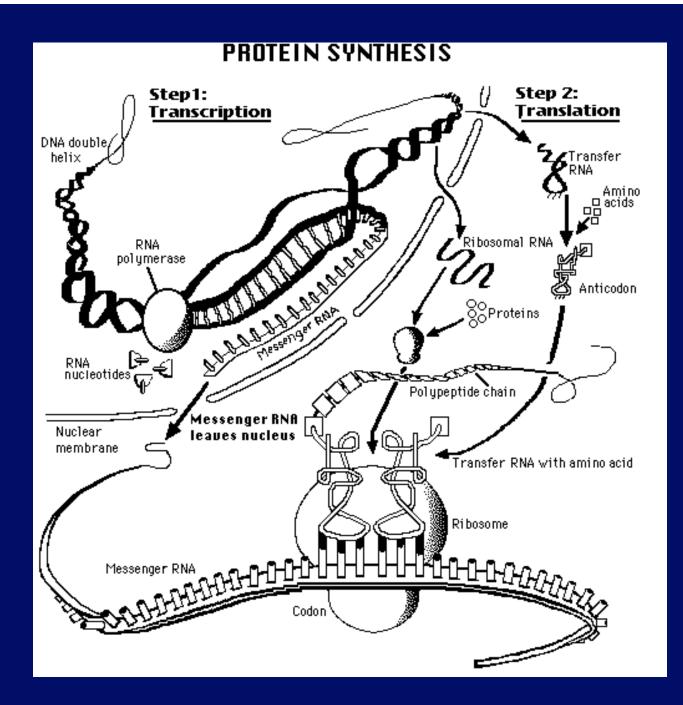
#### Eigen scenario

- A replicating RNA molecule forms by chance (random replicator - not a gene) ribozyme (catalyst, made of RNA)
- 2. Family of **similar** RNA's develops (quasispecies)
- 3. Connection to proteins (quasispecies specialize to make parts of protein)

- 4. Complex interactions (cycles) between RNA and proteins
- 5. Use lipids to make protocells
- 6. Competition leads to biological evolution

### The Origin of the Genetic Code

- We need more than either protein or RNA protolife
- Need interaction via genetic code
- Need translation
- Let's recall what is needed for translation...



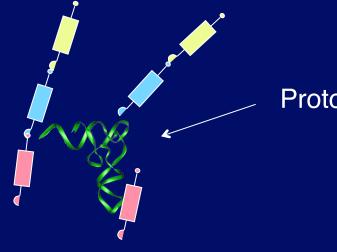
# Shapiro's Fable

The case for the "chicken" Protein first ⇒ replication problem "interpreters" aminoacyl tRNA synthetases

Match tRNA &

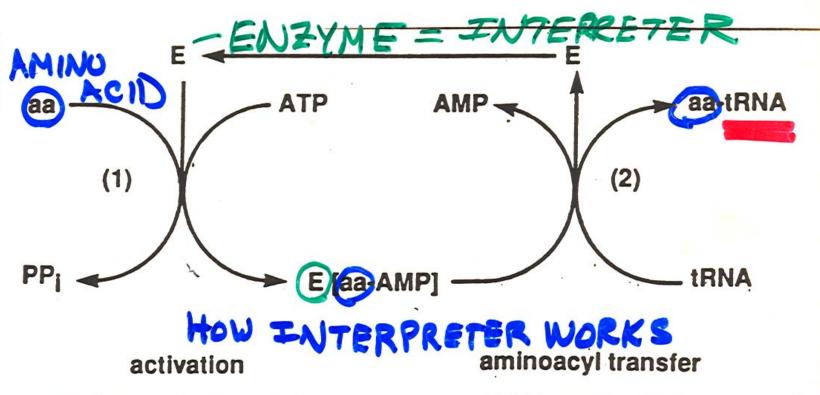
Amino acids

Could an earlier version have copied proteins directly?



**Proto-interpreter** 

tRNA



amino acid recognition

tRNA recognition

Fig. 1 The two steps of the reaction catalysed by aminoacyl-tRNA synthetases. The tRNA is recognized in the second step (through the features designated paracodon in this article) by the enzyme carrying a bound aminoacyl-AMP intermediate. Participation of the aminoacyl group in the recognition process is thus an attractive possibility.

- Early Evolution: Start with 4-6 amino acid types, gradually add more enzymes increase in size and catalytic power
- 2. First use of phosphate as energy? (ATP) or sugar-phosphate chains for construction (Teichoic acids in membranes of some bacteria) (partial  $Q_{\beta}$  replicase)
- Bases added for structure
   Support for protein synthesis → ribosome

- 4. Begin to copy RNA (Full  $Q_{\beta}$  replicase) Natural selection leads to better ribosome
- 5. Specialized, short RNA aided attachment of amino acids to proteins; became tRNA
- Then mRNA to align tRNA's now a separate genetic system that evolves
- 7. DNA developed from RNA

Shapiro dates last step to prokaryote -eukaryote split (different ways of storing DNA info)

Tests:

- 1. Synthesize in lab? Not possible yet.
- 2. Molecular archaeology vestigial ability of interpreters to recognize amino acids in proteins
- 3. Survivors of protein era? prions?

#### Support for the "chicken"

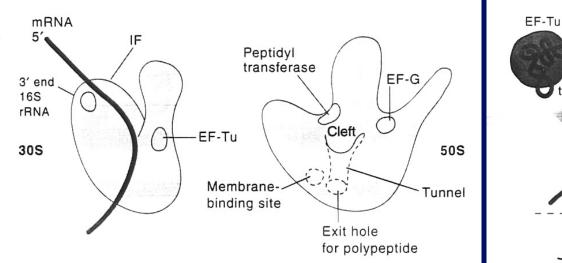
- 1. 1988 discovery that interpreter does not use tRNA codon to recognize correct tRNA (in some cases) ~ 1/2
  - instead a single base pair at the other end of tRNA
  - ⇒ simpler, older code second genetic code
  - ⇒ connection of interpreter and tRNA more primitive than current code

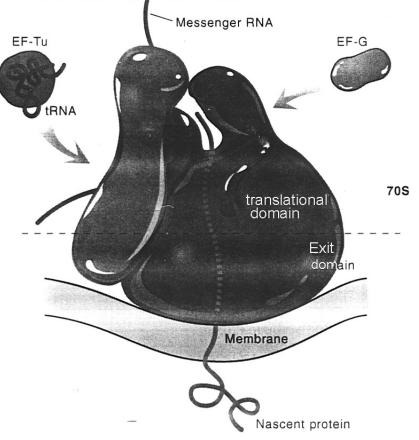
### The Egg Strikes Back

Other work shows some RNA can catalyze Non-RNA reactions

- 1. RNA in ribosome appears to be what catalyzes peptide bond formation Noller, et al. 1992, *Science*, **256**, 1416
- RNA "ribozyme" catalyzes reactions between amino acids and tRNAs
   First "interpreter" may have been RNA Piccirilli, et al. 1992, *Science*, **256**, 1420

## Translation





### New Work

- Progress in RNA copying (Szostak)
- Need Mg<sup>+2</sup> ions, but they can be destructive
- Add citrate, avoids harmful effects of Mg<sup>+2</sup> while keeping good effects
- Need a "leaky" lipid membrane to enclose, but let building blocks in, but keep RNA inside
- Need RNA and lipids
- Science Nov 29, 2013, News Focus

### **Origin of the Genetic Code**

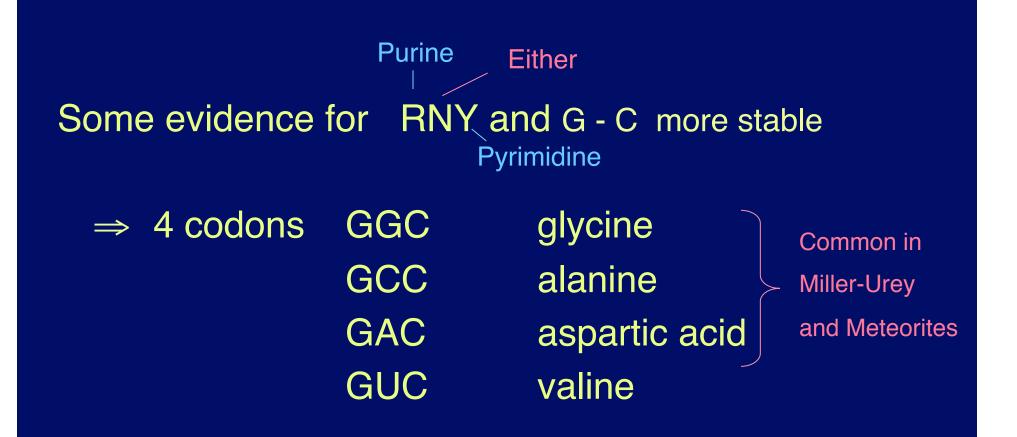
Crucial step in any theory

Allows communication Nucleic Acids ←→→ Proteins

Early versions probably coded fewer amino acids - less specific

For mRNA		Genetic Code			
First RNA Base	U	С	A	G	Third RNA BASE
	Phenylalanine	Serine	Tyrosine	Cysteine	U
	Phenylalanine	Serine	Tyrosine	Cysteine	С
U	Leucine	Serine	Stop	Stop	А
	Leucine	Serine	Stop	Tryptophan	G
	Leucine	Proline	Histidine	Arginine	U
	Leucine	Proline	Histidine	Arginine	С
С	Leucine	Proline	Glutamine	Arginine	А
	Leucine	Proline	Glutamine	Arginine	G
	Isoleucine	Threonine	Asparagine	Serine	U
	Isoleucine	Threonine	Asparagine	Serine	С
A	Isoleucine	Threonine	Lysine	Arginine	А
	Start/Methionine	Threonine	Lysine	Arginine	G
	Valine	Alanine	Aspartic Acid	Glycine	U
	Valine	Alanine	Aspartic Acid	Glycine	С
G	Valine	Alanine	Glutamic Acid	Glycine	А
	Valine	Alanine	Glutamic Acid	Glycine	G



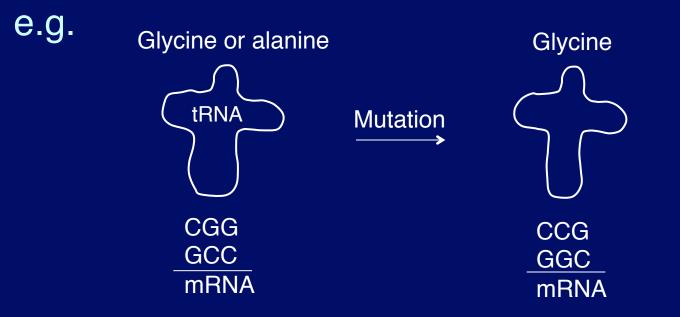


#### Others added later

### **Evolution of Genetic Code**

Gaining specificity

If early tRNAs carried more than 1 kind of amino acid



#### Evidence that code has evolved Freeland, et al. Tested 10<sup>6</sup> other codes

Only one better at minimizing bad effects of mutations

⇒ Natural Selection

Still Evolving Some organisms have slightly different codes in mitochondria or in nucleus

## Summary

- Transition to life is poorly understood
- Need to consider "protolife"
- Can we get by with only one polymer?
  - If so, protein or RNA?
  - If so, how do we get genetic code going?
  - Translation is key
- Most work now focused on nucleic acids
  - Ribozymes can play role of proteins