The Transition to Life

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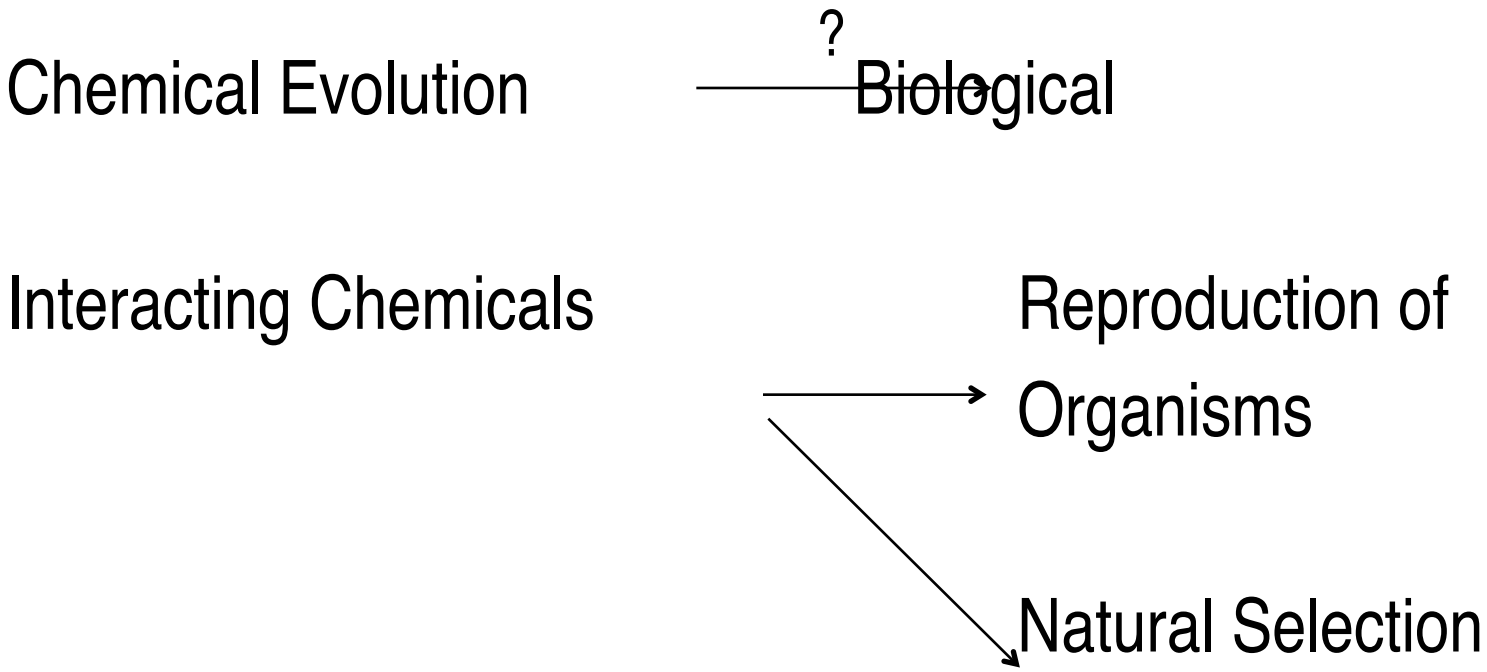


Based on Simplest Life Now:

Need:

1.	Nucleic Acids	Replicable Information
2.	Proteins	Enzymes (Catalysts)
3.	Lipids	Membranes (Enclosure)
4.	Carbohydrates (Pigments)	Energy Storage (Energy Conversion)

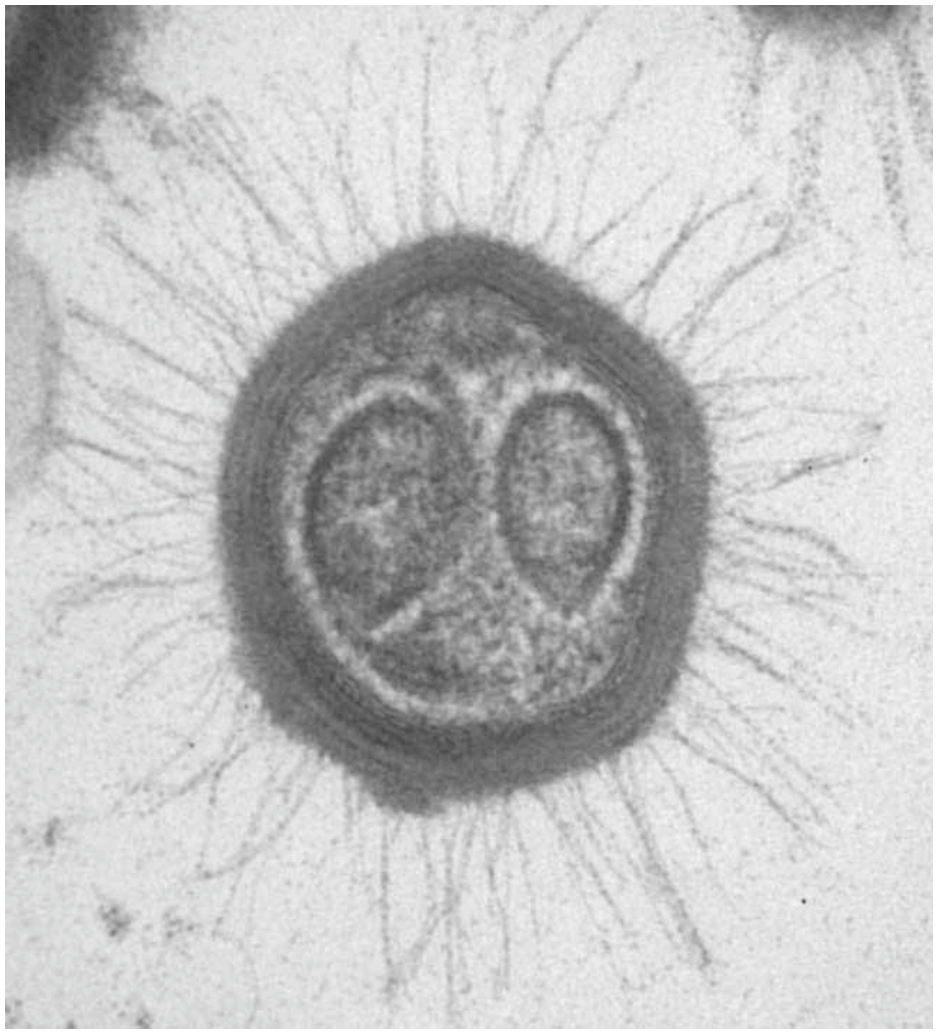
Too much to ask of chemical evolution

$\Rightarrow$ Protolife?

Update: Mimivirus

- A very large virus was discovered in 2003
- Both RNA and DNA
- More DNA than some bacteria
- Genes for translation, DNA repair enzymes
- Leading to reevaluation of viruses
- May be ancient lineages
  - Precursors to bacteria, etc.
  - Controversial

Image of Mimivirus



Protolife

1. “Virus” Free living but equivalent in complexity

   Protein + Nucleic Acid + Supply by Environment

   Genetic Code

2. Protein Protolife

   Protein ~~Self~~ Replication?

3. Nucleic Acid Protolife

   RNA $\longrightarrow$ Self Catalysis?

4. Something Else

   Minerals

   Clay Layers

   Mineral - Molecule

   Pyrite

   Thioesters

Genetic Takeover

? $\longrightarrow$ RNA $\longrightarrow$ DNA

Protein-Based Protolife

1. Proteinoid microspheres - Sidney Fox

   Amino Acids + Dry Heat → Proteinoids

   (Hot Tidepool?) ↓ H_2O (Tide)

   Microspheres

Protocells

Protolife?

Can Add Proteinoid

- Split
- Bud
- Form Chains

(Look like life)

Grow

Divide } "Reproduce"

Bud

Like Bacteria

But "Reproduction" not exact

Later incorporate Nucleic Acids

Proteinoid → Cells → Genes

<u>Problem:</u> How to incorporate Nucleic acids?

Picture of Proteinoid Microspheres

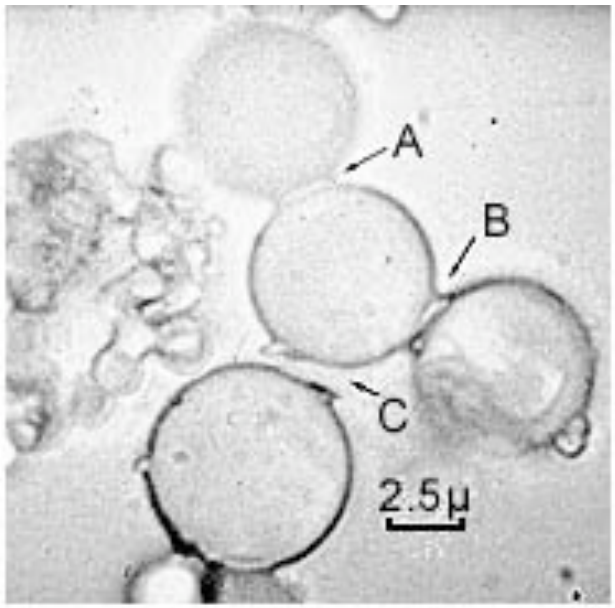


FIGURE 5.15 — Photograph of proteinoid microspheres produced by repeated energizing and dehydrating the primordial soup. The main features of this figure can be simulated by shaking a mixture of oil and water and watching the globs of oil cluster on the surface of the water. Seen here through a microscope, 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~~ ~~Cells~~

Self-replicating RNA molecules

Experiment by Sol Spiegelman

RNA from Q_β 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)

$\Rightarrow$ 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

1. 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 (hypercycles)

6. Use lipids to make protocells

6. Competition leads to biological evolution

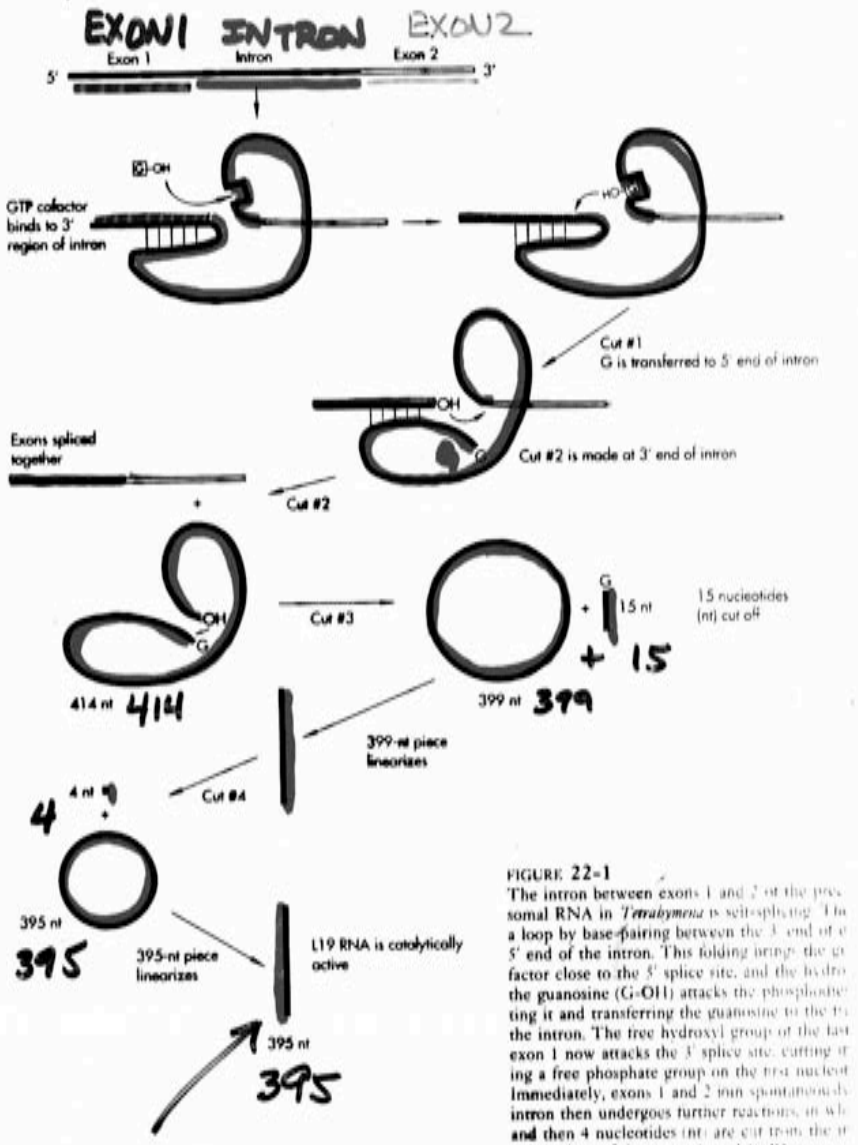


FIGURE 22-1
The intron between exons 1 and 2 of the pre-
somal RNA in *Tetrahymena* is self-splicing. Thi
a loop by base-pairing between the 3' end of e
5' end of the intron. This folding brings the gu
factor close to the 5' splice site, and the hydro
the guanosine (G-OH) attacks the phosphodiest
ting it and transferring the guanosine to the 5'
the intron. The free hydroxyl group of the last
exon 1 now attacks the 3' splice site, cutting it
ing a free phosphate group on the first nucleot
Immediately, exons 1 and 2 join spontaneously
intron then undergoes further reactions, in wh
and then 4 nucleotides (nt) are cut from the 3'
nal product of these reactions, L19 RNA (sinc
19 nucleotides have been removed) is catalytic
It can act as both a polymerase and a nuclease
on pH and the concentrations of the substrates

HOW AN "INTRON" (NOT A GENE) CAN CUT, SPLICE + BECOME A CATALYST

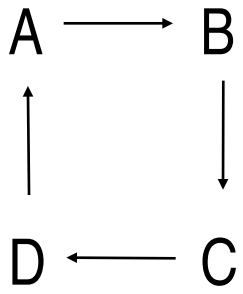
Problems with Nucleic Acid First Scenario

1. Hard to get monomers
2. Unlikely to link correctly
3. Need existing proteins and lipids
4. Hypercycles subject to instabilities

N = size of molecular population

If N small → Population Collapse

If N large → Selfish RNA, Short Circuit



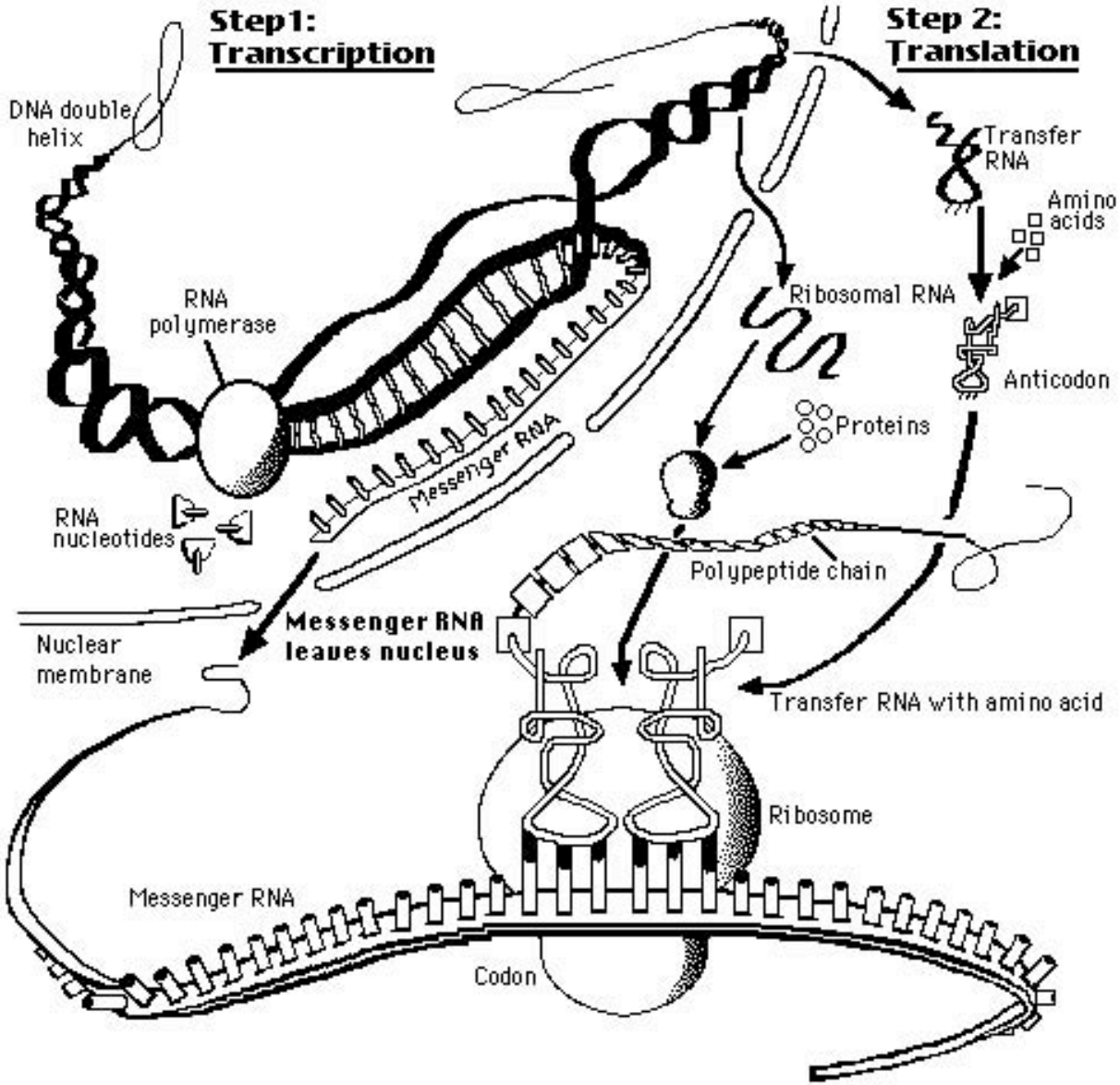
If B → D Short Circuit

$\Rightarrow$ Only narrow range of sizes works

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…

PROTEIN SYNTHESIS

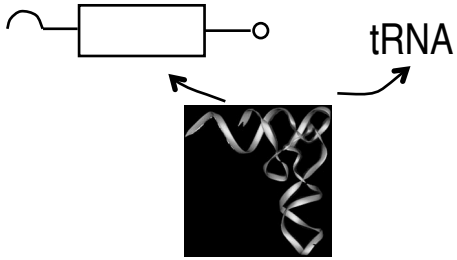


Shapiro's Fable

The case for the "chicken"

Protein first ⇒ replication problem

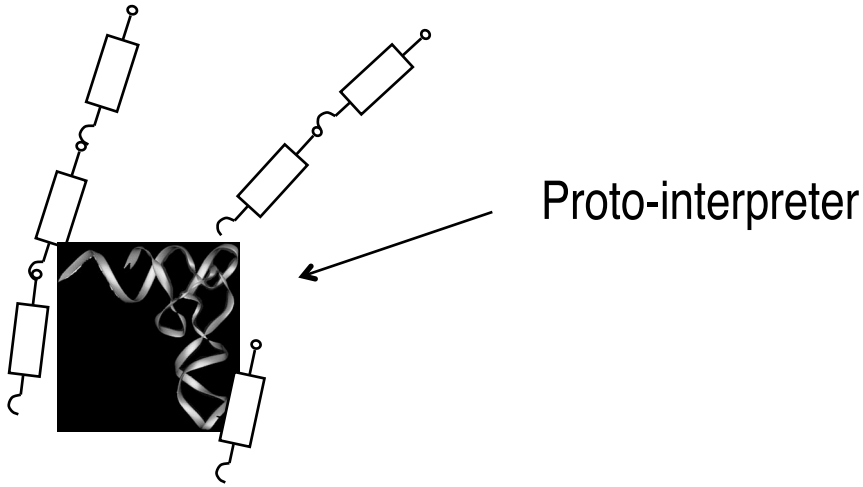
"interpreters" aminoacyl tRNA synthetases



Match tRNA &

Amino acids

Could an earlier version have copied proteins directly?



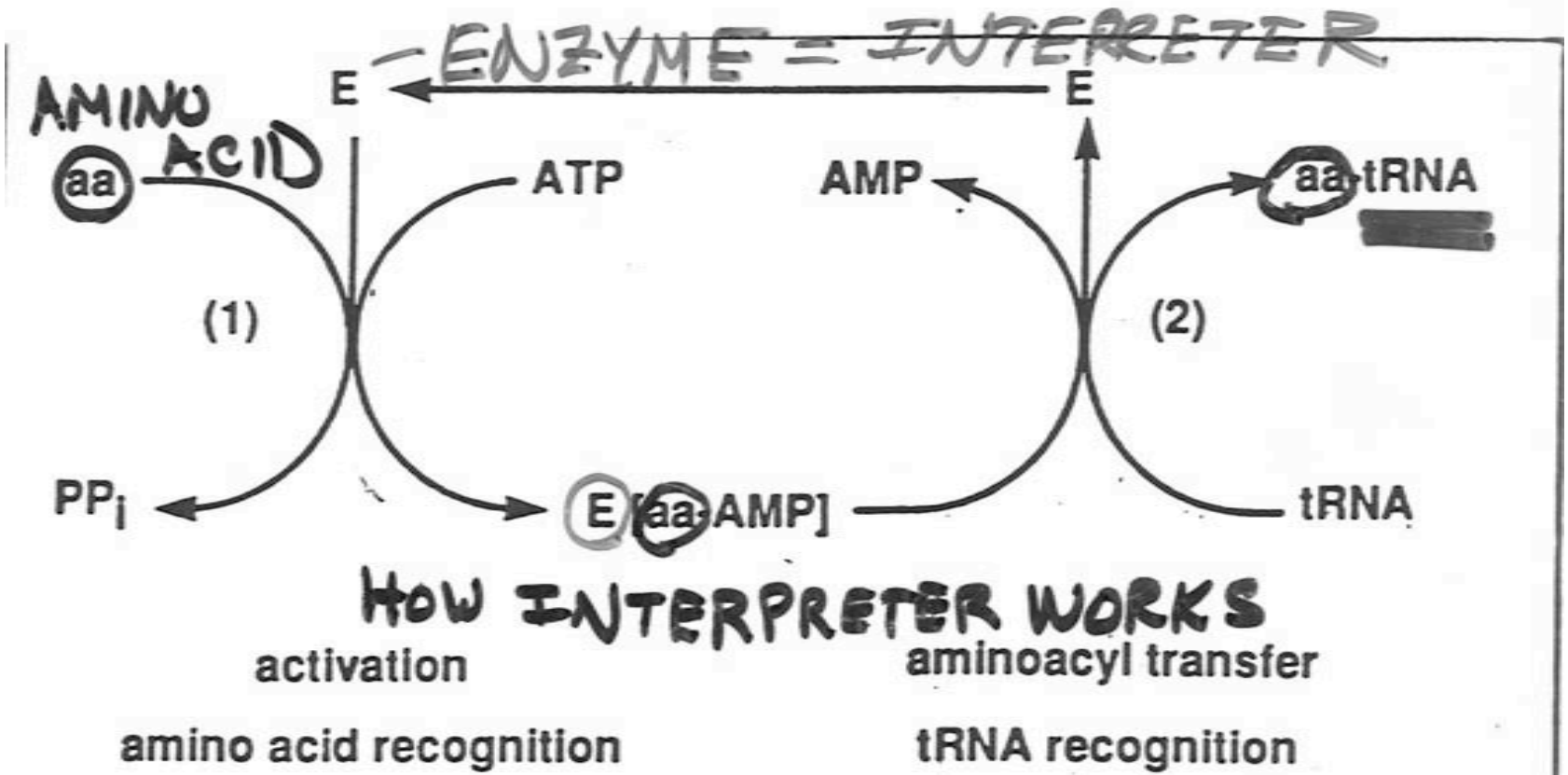


Fig. 1 The two steps of the reaction catalysed by aminoacyl-tRNA synthetases. The tRNA is recognized in the second step (through the features designated paracodon in this article) by the enzyme carrying a bound aminoacyl-AMP intermediate. Participation of the aminoacyl group in the recognition process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

1. 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_β replicase)

3. Bases added for structure
   Support for protein synthesis ~~ribosome~~

4. Begin to copy RNA (Full Q_β replicase)
   Natural selection leads to better ribosome

5. Specialized, Short RNA aided attachment of
   amino acids to proteins; became tRNA

6. 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.

3. 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
   $\Rightarrow$ simpler, older code
   second genetic code

   $\Rightarrow$ connection of interpreter and tRNA
   more primitive than current code

2. Dyson modeling of molecular “populations”

Transition from disorder to order

(non-life) (life)

Finds number of monomer types likely to be

9 - 11 (ok if used ~ 1/2 of modern proteins)

But nucleotides (only 4) - not enough

Favors protein first

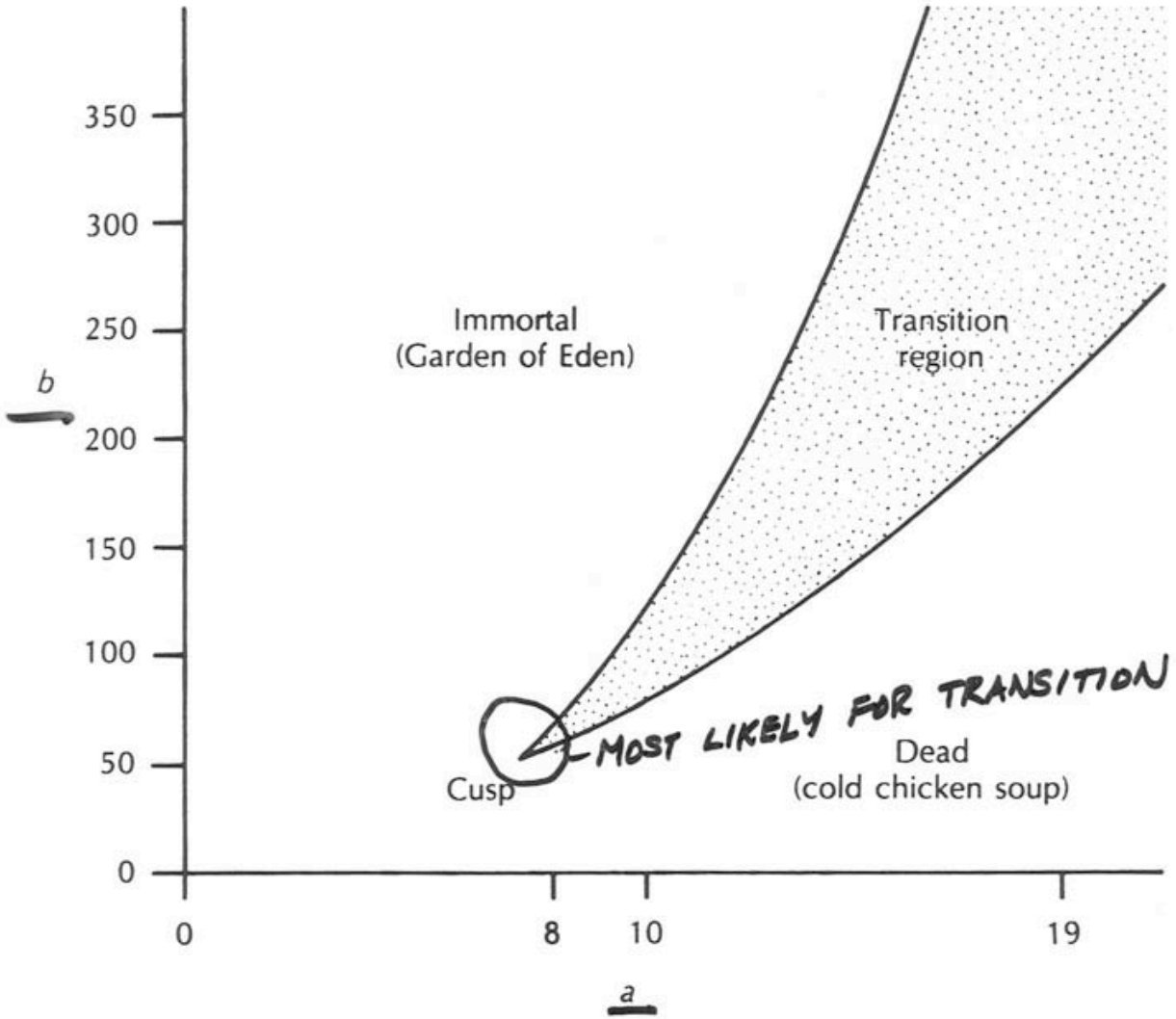


FIGURE 2.7 *Summary of Dyson's model*

1+a = # OF TYPES OF MONOMERS

b = DISCRIMINATION FACTOR OF CATALYST

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

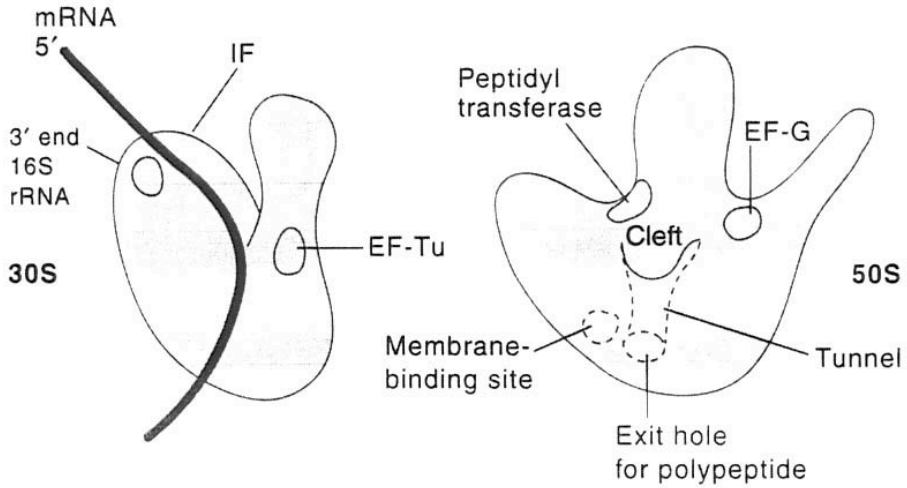
   Noller, et al. 1992, *Science*, **256**, 1416

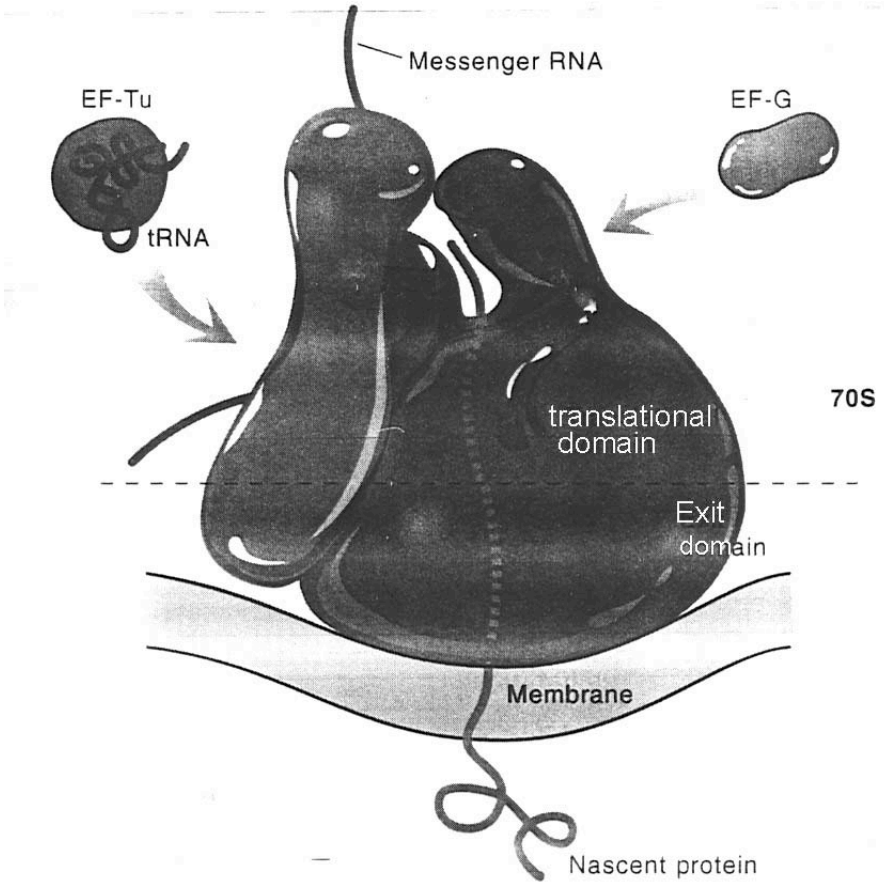
2. RNA “ribozyme” catalyzes reactions between amino acids and tRNAs

   First “interpreter” may have been RNA

   Piccirilli, et al. 1992, *Science*, **256**, 1420

Translation





Origin of the Genetic Code

Crucial step in any theory

Allows communication

Nucleic Acids $\longleftrightarrow$ Proteins

Early versions probably coded fewer amino acids - less specific

For mRNA Genetic Code

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

Amino Acids

Some evidence for RNY and G - C more stable

Purine

Either

Pyrimidine

$\Rightarrow$ 4 codons

GGC	glycine
GCC	alanine
GAC	aspartic acid
GUC	valine

Common in Miller-Urey and Meteorites

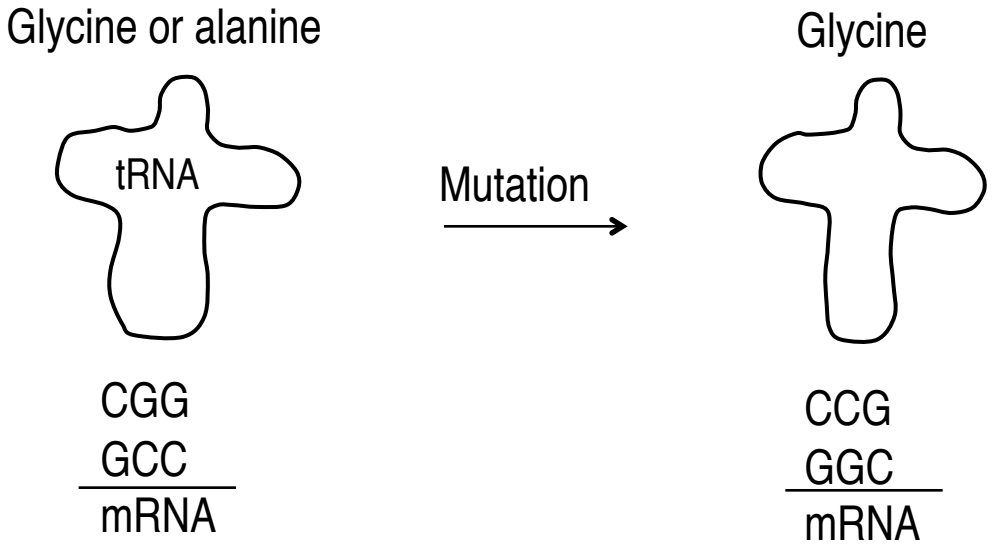
Others added later

Evolution of Genetic Code

Gaining specificity

If early tRNAs carried more than 1 kind of amino acid

e.g.



Evidence that code has evolved
Freeland, et al. Tested 10^6 other codes

Only one better at minimizing bad effects of mutations

$\Rightarrow$ 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?