



Lipids

Evolving chemical systems require compartments for Darwinian evolution – to compete, to store information and to concentrate reactants/metabolites



Encapsulation into membranes is considered an early stage in prebiotic chemical evolution and essential requirement for the emergence of life

Formation of membranes is most easy to explain among major cellular components of the prebiotic Earth.

Many amphiphilic organic compounds spontaneously form vesicles in water at sufficiently high concentrations



Levine, R.M., Pearce, T.R., Adil, M., Kokkoli, E. Langmuir, 2013, 29 (29): 9208–9215.

The vesicle will encapsulate an aqueous solution inside a thin layer of organic material

Modern biological membranes consist primarly of phospholipids with embedded transmembrane proteins. Characterized by low permeability – a disadvantage during early evolution



Fatty acids and fatty alcohols are likely prebiotic lipids

Fischer-Tropsch synthesis

Long hydrocarbon chains from CO + H₂ in presence of metal catalysts and high pressure, fatty acids and alcohols are minor by-products



The mixture of D₂ and CO over meteoritic iron or iron ore produced alkanes and n-fatty acids Oro, J. et al. Geochim. Acta **1976**, 40, 915-924.

Fischer-Tropsch synthesis

Main reactions		
1. Paraffins	$(2n+1)H_2+nCO \rightarrow C_nH_{2n+2}+nH_2O$	
2. Olefins	$2nH_2\text{+}nCO \text{-} C_nH_{2n}\text{+}nH_2O$	
Side reactions		
3. Water-Gas-Shift (WGS)	$CO+H_2O\leftrightarrow CO_2+H_2$	
4. Carbide formation	$yC + xM \leftrightarrow M_xC_y$	
5. Alcohols	$2nH_2+nCO \rightarrow C_nH_{2n}+2O + (n-1)H_2O$	
6. Boudouard reaction	$2CO \rightarrow C + CO_2$	
7. Catalyst reduction and oxidation	$M_xO_y + yH_2 \leftrightarrow xM + yH_2O$	
	$M_xO_y + yCO \leftrightarrow xM + yCO_2$	
8. Coking	$H_2 + CO \rightarrow C + H_2O$	

Hydrothermal Fischer-Tropsch synthesis



Formic or oxalic acid heated in water at 150-250^oC (stainless steel reactor) yielded a mixture of C_{12} - C_{33} lipids

> Rushdi, A., Simoneit B. Origins Life Evol. Biospheres **2001**, *31*, 103-118

When CO, H_2 and NH_3 are allowed to react at 200-700°C in presence of Ni, Al, or clay catalysts, aminoacids are detected:

glycine, alanine, sarcosine, aspartic acid, glutamic acid, arginine, histidine, lysine and ornithine Yoshino, D.; Hayatsu, R.; Anders, E. Geochim. Cosmochim. Acta **1971**, 35, 927-938

Extraterrestrial origin of biomolecules





Murchison meteorite *chondrite*

Table 1. Soluble Organic Compounds in the Murchison Meteorite a

class of compounds	parts per million	n ^b
aliphatic hydrocarbons	>35	140
aromatic hydrocarbons	15-28	87
polar hydrocarbons	<120	10^d
carboxylic acids	>300	48^d
amino acids	60	75^d
imino acids	nd ^c	10
hydroxy acids	15	7
dicarboxylic acids	>30	17^d
dicarboximides	>50	2
pyridinecarboxylic acids	>7	7
sulfonic acids	67	4
phosphonic acids	2	4
N-heterocycles	7	31
amines	13	20^d
amides	nd ^c	27
polyols	30	19

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Fatty acids have been found in meteorites – plausible prebiotic synthesis pathways existed in the early Solar System



Decanoic acid

Extracts of meteorites containing these compounds spontaneously form vescicles when hydrated

Spontaneous generation of lipid vesicles

The first protocell membranes may have assembled from fatty acids and related single-chain lipids available in the prebiotic environment.



At different concentrations, fatty acids can partition between several different phases, including soluble monomers, micelles, and lamellar vesicles, with higher concentrations favoring larger vesicle aggregates.

concentration

I. Budin, A. Debnath, J. W. Szostak J. Am. Chem. Soc., 2012, 134, 20812-20819





By Markus Pössel (Mapos)

Jack Szostak

(* November 9, 1952) - Canadian American biologist of Polish British descent,

Nobel Prize laureate 2009 for Physiology and Medicine, for the discovery of how chromosomes are protected by telomeres; Professor of Genetics at Harvard Medical School.

Szostak has made significant contributions to the field of genetics. His achievement helped scientists to map the location of genes in mammals and to develop techniques for manipulating genes.

His research findings in this area are also instrumental to the Human Genome Project.

In the early 90s his laboratory shifted its research direction and focused on studying **RNA enzymes**, which had been recently discovered by Cech and Altman. He developed the technique of *in vitro evolution of RNA* (also developed independently by Gerald Joyce) which enables the discovery of RNAs with desired functions through successive cycles of selection, amplification and mutation. He isolated the first *aptamer* (term he used for the first time). He isolated *RNA enzymes with RNA ligase activity* directly from random sequence (project of David Bartel).

Currently his lab focuses on the challenges of understanding the *origin of life* on Earth, and the construction of *artificial cellular life* in the laboratory

Coupled growth and division of model protocell membranes



The growth of large multilamellar fatty acid vesicles fed with fatty acid micelles, in a solution where solute permeation across the membranes is slow, results in the transformation of initially spherical vesicles into long thread-like vesicles, a process driven by the transient imbalance between surface area and volume growth. Modest shear forces are then sufficient to cause the thread-like vesicles to divide into multiple daughter vesicles without loss of internal contents.



Ting F. Zhu, and Jack W. Szostak J. Am. Chem. Soc., 2009, 131 (15), 5705-5713

Coupled growth and division of model protocell membranes



Cycles of vesicle growth and division. (A) Relative surface area after two cycles of addition of 5 equiv of oleate micelles (solid circles) or 5 equiv of NaOH (open circles) to oleate vesicles, each followed by agitation. Inset micrographs show vesicle shapes at indicated times. Scale bar, 10 μ m. (B) Vesicle shapes during cycles of growth and division in a model prebiotic buffer (0.2 M Na-glycine, pH 8.5, ~1 mM initial oleic acid, vesicles contain 10 mM HPTS for fluorescence imaging). Scale bar, 20 μ m.

Ting F. Zhu, and Jack W. Szostak J. Am. Chem. Soc., 2009, 131 (15), 5705-5713



B С D

Photochemically driven protocell division

The illumination of filamentous fatty acid vesicles rapidly induces pearling and subsequent division in the presence of thiols.

Photochemically generated reactive oxygen species oxidize thiols to disulfidecontaining compounds that associate with fatty acid membranes, inducing a change in surface tension and causing pearling and subsequent division.

Alternative route for the emergence of early self-replicating cell-like structures, particularly in thiol-rich surface environments. The subsequent evolution of cellular metabolic processes controlling the thiol:disulfide redox state would have enabled autonomous cellular control of the timing of cell division, a major step in the origin of cellular life.

Oleate vesicle pearling and division.

A. Radical-mediated oxidation of DTT.

B. An oleate vesicle (containing 2 mM HPTS, in 0.2 M Na-glycinamide,

pH 8.5, 10 mM DTT) 30 min after the addition of five equivalents of oleate micelles.

C. and D. Under intense illumination (for 2 s and 12 s, respectively), the long thread-like vesicle went through pearling and division Scale bar, $10\mu m$.

T. F. Zhu, K. Adamala, N. Zhang, J. W. Szostak PNAS, 2012, doi:10.1073/pnas.1203212109



Photochemically driven protocell division

Oleate vesicle pearling and division with various thiols in the solution.

(A) 3-mercaptopropionic acid.

(B and C) An oleate vesicle (containing 2 mM HPTS, in 0.2 M Na-bicine, pH 8.5, 10 mM 3-mercaptopropionic acid, 30 min after the addition of five equivalents of oleatemicelles) went through pearling and division under intense illumination (for 3 s and 15 s, respectively).

(D) 3-mercapto-1-propanol.

(E and F) An oleate vesicle as above but in 50 mM 3-mercapto-1-propanol, went through pearling and division under intense illumination (for 2 s and 10 s, respectively).

(G) 1-mercapto-2-propanol.

(H and I) An oleate vesicle as above but in 50 mM 1-mercapto-2-propanol went through pearling and division under intense illumination (for 2 s and 9 s, respectively).

(J) 3-mercapto-1,2,4-triazole.

(K and L) An oleate vesicle as above but in 50 mM 3-mercapto-1,2,4-triazole went through pearling and division under intense illumination (for 3 s and 13 s, respectively). Scale bar, 20 μ m.

T. F. Zhu, K. Adamala, N. Zhang, J. W. Szostak *PNAS*, **2012**, doi:10.1073/pnas.1203212109





RNA localization with a model amphipathic, cationic molecule

) Design of RNA-localizing molecules that include both nonpolar and cationic regions.

b) The change in zeta potential

c) Schematic of the FRET assay used to assess RNA localization to vesicle membranes

d) RNA (5'-FAM-U₁₅ and 5'-FAM-A₁₅) shows increasing localization to POPC membranes that contain increased amounts of undecylimidazole.





Microscopy of encapsulated RNA localization to POPC membranes with 2-undecylimidazole. Confocal images of 5'-FAM-U₁₅RNA (green) association with giant POPC vesicles membranes in the presence of 2-undecylimidazole. Differential interference contrast (DIC) microscopy images are shown for each vesicle.

- a) RNA appears uniformly distributed in the interior of POPC GUVs.
- b) The addition of SUVs containing a rhodamine-labeled lipid (red) leads to SUV aggregation and association with the giant vesicle membranes, but RNA (green) remains uniformly encapsulated in the vesicle interior.
- c) The addition of SUVs containing a rhodamine-labeled lipid (red) and 40 mol% 2-undecylimidazole leads to SUV association with vesicle membranes and RNA (green) localizes to the vesicle surface. The scale bar is 20 mm.

SUV – small unilamellar vescile GUV – giant unilamellar vescile (5-25 μ m)



Peptide-induced RNA-membrane association. A FRET assay reports RNA localization (5'-FAM-U₁₅) to POPC and oleic acid membranes (7.5 mm) 10 h after the addition of 1 mm of various peptides to the vesicle solution at pH 8. Data is reported as a percentage change from control samples that lack peptide. n=4, error bars represent the standard error of the mean.



Microscopy of peptide-induced RNA–membrane association. Confocal images show RNA localization (5'-AlexaFluor647-labeled 15-mer, cyan) to the outside of oleic acid/POPC (90%/10%) and pure POPC membranes in the presence of R3F3 and R3W3 peptides. Control samples had no peptide added. For each image, the left panel shows the DIC image and the right panel shows AlexaFluor647 fluorescence. The scale bar is 20 mm.



Microscopy of encapsulated RNA localization to POPC membranes with peptides. Confocal images show that RNA (5'-FAMU₁₅, green) encapsulated in POPC vesicles (containing a rhodaminelabeled lipid, red) becomes localized to the membrane of certain vesicles after an overnight incubation with R3F3 and R3W3 peptides. The scale bar is 20 mm.

Phosphates



Schreibersite is generally a rare iron-nickel phosphide mineral, $(Fe,Ni)_3P$, though common in iron-nickel meteorites Acidic schreibersite corrosion under anaerobic conditions $(10\% \text{ aq. HCl/N}_2) \rightarrow \text{ soluble forms of phosphorus}$

 $(Fe,Ni)_{3}P + HCl_{aq} \rightarrow H_{2}PO_{3}^{-} \rightarrow H_{2}P_{2}O_{5}^{2-}$

T. P. Kee *et al.Geochimica et Cosmochimica Acta*. **2013** *109,* 90-112



slice of the Gebel Kamil Meteorite with schreibersite rimmed by kamacite



Virginia Smith, UA Lunar & Planetary Laboratory

Image of schreibersite grain present in a thinsection of the enstatite meteorite, KLE 98300.

Phosphates



organophosphorous compounds are detected

I. Acetyl-P compounds $= C_2 O_2 H_3$ C2O2H3 HO HO C2O2H3 HO C2O2H2 HO Phosphonoacetate Phosphatoacetate Hypophosphatoacetate HO HO-C₂O₂H₃ HO HO C2O2H3 Etc. Tri-P-acetate Quad-P-acetate Penta-P-Acetate II. Methyl-P Compounds OH OH HO OH H_2 Phosphonomethanol Dimethylphosphite Phosphonomethylphosphate III. Phosphonoformate HO-OH. но

M. Pasek et al. Geochimica et Cosmochimica Acta. 2007 71, 1721-1736

Phospholipids



M. Powner, J. Sutherland Phil. Trans. R. Soc. B 2011, 366, 2870–2877

Lipids - summary

Many amphiphilic organic compounds spontaneously form vesicles in water at sufficiently high concentrations

Current phospholipid membranes likely evolved late. Protocells probably encapsulated by fatty acids, fatty alcohols, prenyl oligomers, or phosphorylated alcohols

Nucleolipids are proposed as intermediates in templated oligonucleotide replication

Phosphorus was accessible upon corrosion of meteorite materials and could be incorporated into lipids



The origin of small reactive intermediates

Schreibersite $(Fe,Ni)_3P$, from iron-nickel meteorites: source of phosphorus, iron and nickel Under more neutral conditions phosphates recombine with iron \rightarrow Fe₃(PO₄)₂ (**vivianite**)

It should be re-solubilized to become accessible for following chemical transformations

HCN – the crucial reactive intermediate – burning of carbon-rich chondrite meteorites into redox-neutral atmosphere containing N₂ and water

 $Fe_3(PO_4)_2 + 18CN_{aq} \rightarrow 2PO_4^{3-} + 3[Fe(CN)_6]^{4-}$

Two important functions: solubilization of phosphates and concentration of atmospheric HCN deposited as salts of monoand divalent cations (Na, K, Mg, Ca)

Similar reactions take place with insoluble copper and nickel sulfides deposited by iron-nickel meteorite impacts (same occurrence as schreibersite, rich mining sources of these metals until today)

NiS + H₂O + 6CN⁻ → [Ni(CN)₆]⁴⁻ + HS⁻ + OH⁻ Cu₂S + H₂O + 6CN⁻ → 2[Cu(CN)₃]²⁻ + HS⁻ + OH⁻

The origin of small reactive intermediates

 $Cu_2S + H_2O + 6CN^2 \rightarrow 2[Cu(CN)_3]^{2-} + HS^2 + OH^2$

Thermal decomposition of cyanoferrates (volcanic):



Action of water (bufferred to neutral or slightly acidic) on that mixture produced concentrated HCN solution + cyanamide (from CaNCN) + acetylene (from CaC₂) + ammonia (from Mg₃N₂)