Cosmology and the Origin of Life*

John D. Barrow
Astronomy Centre
University of Sussex
Brighton BN1 9QJ
U.K.

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Abstract

We discuss some of the cosmological constraints on the evolution and persistence of life in the Universe and in hypothetical universes other than our own. We highlight the role played by the age and size of the universe, and discuss the interplay between the main-sequence stellar lifetime and the biological evolution time scale. The consequences of different versions of the inflationary universe scenario are described in the light of limits on the possible variation in the values of the constants of Nature.

1 Cosmology, Stars and Life

Prior to the discovery of the expansion of the Universe there was little that cosmology could contribute to the question of extraterrestrial life aside from probabilities and prejudices. After our discovery of the expansion and evolution of the Universe the situation changed significantly. The entire cosmic environment was recognised as undergoing steady change. The history of the Universe took on the complexion of an unfolding drama in many acts, with

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the formations of first atoms and molecules, then galaxies and stars, and most recently, planets and life. The most important and simplest feature of the overall change in the Universe that the expansion produces is the rate at which it occurs. This is linked to the age of the expanding universe and that of its constituents.

In the 1930s, the distinguished biologist JBS Haldane took an interest in Milne's proposal \[1\] that there might exist two different timescales governing the rates of change of physical processes in the Universe: one, \(t\), for 'atomic' changes and another, \(\tau\), for 'gravitational changes' where \(\tau = \ln(t/t_0)\) with \(t_0\) constant. Haldane explored how changing from one timescale to the other could alter one's picture of when conditions in the Universe would become suitable for the evolution of biochemical life \[2\], \[4\]. In particular, he argued that it would be possible for radioactive decays to occur with a decay rate that was constant on the \(t\) timescale but which grew in proportion to \(t\) when evaluated on the \(\tau\) scale. The biochemical processes associated with energy derived from the breakdown of adenosine triphosphoric acid would yield energies which, while constant on the \(t\) scale, would grow as \(t^2\) on the \(\tau\) scale. Thus there would be an epoch of cosmic history on the \(\tau\) scale before which life was impossible but after which it would become increasingly likely. Milne's theory subsequently fell into abeyance although the interest in gravitation theories with a varying Newtonian 'constant' of gravitation led to detailed scrutiny of the paleontological and biological consequences of such hypothetical changes for the past history of the Earth \[4\]. Ultimately, this led to the formulation of the collection of ideas now known as the Anthropic Principles, \[5\], \[6\].

Another interface between the problem of the origin of life and cosmology has been the perennial problem of dealing with finite probabilities in situations where an infinite number of potential trials seem to be available. For example, in a universe that is infinite in spatial volume (as would be expected for the case for an expanding open universe with non-compact topology), any event that has a finite probability of occurring should occur not just once but infinitely often with probability one if the spatial structure of the Universe is exhaustively random \[3\]. In particular, in an infinite universe we conclude that there should exist an infinite number of sites where life has progressed to our stage of development. In the case of the steady-state universe, it is possible to apply this type of argument to the history of the universe as well as its geography because the universe is assumed to be infinitely old. Every past-directed world line should encounter a living civilisation. Accordingly,
it has been argued that the steady state universe makes the awkward prediction that the universe should now be teeming with life along every line of sight [4].

The key ingredient that modern cosmology introduces into considerations of biology is that of time. The observable universe is expanding and not in a steady state. The density and temperature are steadily falling as the expansion proceeds. This means that the average ambient conditions in the universe are linked to its age. Roughly, in all expanding universes, dimensional analysis tells us that the density of matter, \( \rho \), is related to the age \( t \) measured in comoving proper time and Newton’s gravitation constant, \( G \), by means of a relation of the form

\[
\rho \approx \frac{1}{Gt^2}
\]  

(1)

The expanding universe creates an interval of cosmic history during which biochemical observers, like ourselves, can expect to be examining the Universe. Chemical complexity requires basic atomic building blocks which are heavier than the elements of hydrogen and helium which emerge from the hot early stages of the universe. Heavier elements, like carbon, nitrogen, and oxygen, are made in the stars, as a result of nuclear reactions that take billions of years to complete. Then, they are dispersed through space by supernovae after which they find their way into grains, planets, and ultimately, into people. This process takes billions of years to complete and allows the expansion to produce a universe that is billions of light years in size. Thus we see why it is inevitable that the universe is seen to be so large. A universe that is billions of years old and hence billions of light years in size is a necessary pre-requisite for observers based upon chemical complexity. Biochemists believe that chemical life of this sort, and the form based upon carbon in particular, is likely to be the only sort able to evolve spontaneously. Other forms of living complexity (for example that being sought by means of silicon physics) almost certainly can exist but it is being developed with carbon-based life-forms as a catalyst rather than by spontaneous evolution.

The inevitability of universes that are big and old as habitats for life also leads us to conclude that they must be rather cold on average because significant expansion to large size reduces the average temperature inversely in proportion to the size of the universe. They must also be sparse, with a low average density of matter and large distances between different stars and galaxies. This low temperature and density also ensures that the sky
is dark at night (the so called 'Olbers’ Paradox’ first noted by Halley, [7]) because there is too little energy available in space to provide significant apparent luminosity from all the stars. We conclude that many aspects of our Universe which, superficially, appear hostile to the evolution of life are necessary prerequisites for the existence of any form of biological complexity in the Universe.

Life needs to evolve on a timescale that is intermediate between the typical time scale that it takes for stars to reach a stable state of stable hydrogen burning, the so called main-sequence lifetime, and the timescale on which stars exhaust their nuclear fuel and gravitationally collapse. This timescale, $t_*$, is determined by a combination of fundamental constants of Nature

$$t_* \approx \left( \frac{G m_N^2}{h c} \right)^{-1} \times \frac{h}{m_N c^2} \approx 10^9 \text{ yrs}$$

(2)

where $m_N$ is the proton mass, $h$ is Planck’s constant, and $c$ is the velocity of light [8], [4].

In expanding universes of the Big Bang type the reciprocal of the observed expansion rate of the universe, Hubble’s constant $H_0 \approx 70 \text{Km.s}^{-1} \text{Mpc}^{-1}$, is closely related to the expansion age of the universe, $t_0$, by a relation of the form

$$t_0 \approx \frac{2}{3H_0}$$

(3)

The fact that the age $t_0 \approx 10^{10} \text{ yr}$ deduced from observations of $H_0$ in this way is a little larger than the main sequence lifetime, $t_*$, is entirely natural in the Big Bang theory that is, we observe a little later than the time when the Sun forms). However, the now defunct steady state theory, in which there is no relation between the age of the universe (which is infinite) and the measured value of $H_0$, would have had to regard the closeness in value of $H_0^{-1}$ and $t_*$ as a complete coincidence [9].

## 2 Biology and Stars: Is there a link?

Evidently, in our solar system life first evolved quite soon after the formation of a hospitable terrestrial environment. Suppose the typical time that it takes for life to evolve is denoted by some timescale $t_{bio}$, then from the evidence
presented by the solar system, which is about $4.6 \times 10^9\text{yrs}$ old, it is seems that

\[ t_* \approx t_{bio} \]

At first sight we might assume that the microscopic biochemical processes and local environmental conditions that combine to determine the magnitude of $t_{bio}$ are independent of the nuclear astrophysical and gravitational processes that determine the typical stellar main sequence lifetime $t_{ms}$. However, this assumption leads to the striking conclusion that we should expect extraterrestrial forms of life to be exceptionally rare [10], [4], [11]. The argument, in its simplest form, is as follows. If $t_{bio}$ and $t_*$ are independent then the time that life takes to arise is random with respect to the stellar timescale $t_*$. Thus it is most likely that either $t_{bio} \gg t_*$ or that $t_{bio} \ll t_*$. Now if $t_{bio} \ll t_*$ we must ask why it is that the first observed inhabited solar system (that is, us) has $t_{bio} \approx t_*$. This would be extraordinarily unlikely. On the other hand, if $t_{bio} \gg t_*$ then the first observed inhabited solar system (us) is most likely to have $t_{bio} \approx t_*$ since systems with $t_{bio} \gg t_*$ have yet to evolve. Thus we are a rarity, one of the first living systems to arrive on the scene. Generally, we are led to a conclusion, an extremely pessimistic one for the SETI enterprise, that $t_{bio} \gg t_*$. 

In order to escape from this conclusion we have to undermine one of the assumptions underlying the argument that leads to it. For example, if we suppose that $t_{bio}$ is no independent of $t_*$ then things look different. If $t_{bio}/t_*$ is a rising function of $t_*$ then it is actually likely that we will find $t_{bio} \approx t_*$. Livio [12] has given a simple model of how it could be that $t_{bio}$ and $t_*$ are related by a relation of this general form. He takes a very simple model of the evolution of a life-supporting planetary atmosphere like the Earth's to have two key phases which lead to its oxygen content:

**Phase 1**: Oxygen is released by the photodissociation of water vapour. On Earth this took $2.4 \times 10^9\text{yr}$ and led to an atmospheric $O_2$ build up to about $10^{-3}$ of its present value.

**Phase 2**: Oxygen and ozone levels grow to about 0.1 of their present levels. This is sufficient to shield the Earth's surface from lethal levels of ultra-violet radiation in the 2000-3000 Å band (note that nucleic acid and protein absorption of ultra-violet radiation peaks in the 2600-2700 Å and 2700-2900 Å bands, respectively). On Earth this phase took about $1.6 \times 10^9\text{yr}$. 
Now the length of Phase 1 might be expected to be inversely proportional to the intensity of radiation in the wavelength interval 1000-2000 Å, where the key molecular levels for $H_2O$ absorption lie. Studies of stellar evolution allow us to determine this time interval and provide a rough numerical estimate of the resulting link between the biological evolution time (assuming it to be determined closely by the photodissociation time) and the main sequence stellar lifetime, with \[ t_{bio} \approx 0.4 \left( \frac{t_*}{t_{sun}} \right)^{1.7}, \]

where $t_{sun}$ is the age of the Sun.

This model indicates a possible route to establishing a link between the biochemical timescales for the evolution of life and the astrophysical timescales that determine the time required to create an environment supported by a stable hydrogen burning star. There are obvious weak links in the argument. It provides on a necessary condition for life to evolve, not a sufficient one. We know that there are many other events that need to occur before life can evolve in a planetary system. We could imagine being able to derive an expression for the probability of planet formation around a star. This would involve many other factors which would determine the amount of material available for the formation of solid planets with atmospheres at distances which permit the presence of liquid water and stable surface conditions. Unfortunately, we know that there were many 'accidents' of the planetary formation process in the solar system which have subsequently played a major role in the existence of long-lived stable conditions on Earth, [13]. For example, the presence of resonances between the precession rates of rotating planets and the gravitational perturbations they feel from all other bodies in their solar system can easily produce chaotic evolution of the tilt of a planet’s rotation axis with respect to the orbital plane of the planets over times must shorter than the age of the system [14], [13]. The planet’s surface temperature variations, insolation levels, and sea levels are sensitive to this angle of tilt. It determines the climatic differences between what we call ‘the seasons’. In the case of the Earth, the modest angle of tilt (approximately 23 degrees) would have experienced this erratic evolution had it not been for the presence of the Moon [13], [13]. The Moon is large enough for its gravitational effects to dominate the resonances which occur between the Earth’s precessional rotation and the frequency of external gravitational perturbations from the other planets. As a result the Earth’s tilt wobbles
only by a fraction of a degree around 23° over hundreds of thousands of years. Enough perhaps to cause some climatic change, but not catastrophic for the evolution of life.

This shows how the causal link between stellar lifetimes and biological evolution times may be rather a minor factor in the chain of fortuitous circumstances that must occur if habitable planets are to form and sustain viable conditions for the evolution of life over long periods of time. The problem remains to determine whether the other decisive astronomical factors in planet formation are functionally linked to the surface conditions needed for biochemical processes.

3 Habitable Universes

We know that several of the distinctive features of the large scale structure of the visible universe play a role in meeting the conditions needed for the evolution of biochemical complexity within it.

The first example is the proximity of the expansion dynamics to the ‘critical’ state which separates an ever-expanding future from one of eventual contraction, to better than ten per cent. Universes that expanded far faster than this would be unable to form galaxies and stars and hence the building blocks of biochemistry would be absent. The rapid expansion would prevent islands of material separating out from the global expansion and becoming bound by their own self-gravitation. By contrast, if the expansion rate were far below that characterising the critical rate then the material in the universe would have condensed into dense structures and black holes long before stars could form [16], [17], [1], [19], [18].

The second example is that of the uniformity of the universe. The non-uniformity level on the largest scales is very small, \( \Delta \approx 10^{-5} \). This is a measure of the average relative fluctuations in the gravitational potential on all scales. If \( \Delta \) were significantly larger then galaxies would have rapidly degenerated into dense structures within which planetary orbits would be disrupted by tidal forces and black holes would form rapidly before life-supporting environments could be established. If \( \Delta \) were significantly smaller then the non-uniformities in the density would be gravitationally too feeble to collapse into galaxies and no stars would form. Again, the universe would be bereft of the biochemical building blocks of life [20].

In recent years the most popular theory of the very early evolution of the
universe has provided a possible explanation as to why the universe expands so close to the critical life-supporting divide and why the fluctuation level has the value observed. This theory is called 'inflation'. It proposes that during a short interval of time when the temperature was very high (say $\sim 10^{25} K$), the expansion of the universe accelerated. This requires the material content of the universe to be temporarily dominated by forms of matter which effectively antigravitate for that period of time \[21\]. This requires their density $\rho$, and pressure, $p$, to satisfy the inequality \[19\]

$$\rho + \frac{3p}{c^2} < 0 \quad (4)$$

The inflation is envisaged to end because the matter fields responsible decay into other forms of matter, like radiation, which do not satisfy this inequality. After this occurs the expansion resumes the state of decelerating expansion that it possessed before its inflationary episode began.

If inflation occurs it offers the possibility that the whole of the visible part of the universe (roughly 15 billion light years in extent today) has expanded from a region that was small enough to be causally linked by light signals at the very high temperatures and early times when inflation occurred. If inflation does not occur then the visible universe would have expanded from a region that is far larger than the distance that light can circumnavigate at these early times and so its smoothness today is a mystery. If inflation occurs it will transform the irreducible quantum statistical fluctuations in space into distinctive patterns of fluctuations in the microwave background radiation which future satellite observations will be able to detect if they were of an intensity sufficient to have produced the observed galaxies and clusters by the process of gravitational instability.

As the inflationary universe scenario has been explored in greater depth it has been found to possess a number of unexpected properties which, if they are realised, would considerably increase the complexity of the global cosmological problem and create new perspectives on the existence of life in the universe \[22\], \[23\], \[19\].

It is possible for inflation to occur in different ways in different places in the early universe. The effect is rather like the random expansion of a foam of bubbles. Some inflate considerably while others hardly inflate at all. This is termed 'chaotic inflation'. Of course, we have to find ourselves in one of the regions that underwent sufficient inflation so that the expansion lasted for longer than $t_\star$ and stars could produce biological elements. In
such a scenario the global structure of the Universe is predicted to be highly inhomogeneous. Our observations of the microwave background temperature structure will only be able to tell us whether the region which expanded to encompass out visible part of the universe underwent inflation in its past. An important aspect of this theory is that for the first time it has provided us with a positive reason to expect that the observable universe is not typical of the structure of the universe beyond our visible horizon, 15 billion light years away.

It was subsequently been discovered that under fairly general conditions inflation can be self-reproducing. That is, quantum fluctuations within each inflating bubble will necessarily create conditions for further inflation of microscopic regions to occur. This process or 'eternal inflation' appears to have no end and may not have had a beginning. Thus life will be possible only in bubbles with properties which allow self-organised complexity to evolve and persist.

It has been found that there is further scope for random variations in these chaotic and eternal inflationary scenarios. In the standard picture we have just sketched, properties like the expansion rate and temperature of each inflated bubble can vary randomly from region to region. However, it is also possible for the strengths and number of low-energy forces of Nature to vary. It is even possible for the number of dimensions of space which have expanded to large size to be different from region to region. We know that we cannot produce the known varieties of organised biochemical complexity if the strengths of forces change by relatively small amounts, or in dimensions other than three because of the impossibility of creating chemical or gravitational bound states, [24, 25, 26, 4, 27].

The possibility of these random variations arises because inflation is ended by the decay of some matter field satisfying (4). This corresponds to the field evolving to a minimum in its self-interaction potential. If that potential has a single minimum then the characteristic physics that results from that ground state will be the same everywhere. But if the potential has many minima (for example like a sine function) then each minimum will have different low-energy physics and different parts of the universe can emerge from inflation in different minima and with different effective laws of interaction for elementary particles. In general, we expect the symmetry breaking which chooses the minima in different regions to be independent and random.
4 Changing Constants

Considerations like these, together with the light that superstring theories have shed upon the origins of the constants of Nature, mean that we should assess how narrowly defined the existing constants of Nature need to be in order to permit biochemical complexity to exist in the Universe \[4\], \[28\]. For example, if we were to allow the ratio of the electron and proton masses \((\beta = m_e/m_N)\) and the fine structure constant \(\alpha\) to be change their values (assuming no other aspects of physics is changed by this assumption – which is clearly going to be false!) then the allowed variations are very constraining. Increase \(\beta\) too much and there can be no ordered molecular structures because the small value of \(\beta\) ensures that electrons occupy well-defined positions in the Coulomb field created by the protons in the nucleus; if \(\beta\) exceeds about \(5 \times 10^{-3} \alpha^2\) then there would be no stars; if modern grand unified gauge theories are correct then \(\alpha\) must lie in the narrow range between about \(1/180\) and \(1/85\) in order that protons not decay too rapidly and a fundamental unification of non-gravitational forces can occur. If, instead, we consider the allowed variations in the strength of the strong nuclear force, \(\alpha_s\), and \(\alpha\) then roughly \(\alpha_s < 0.3\alpha^{1/2}\) is required for the stability of biologically useful elements like carbon. If we increase \(\alpha_s\) by 4% there is disaster because the helium-2 isotope can exist (it just fails to be bound by about 70 KeV in practice) and allows very fast direct proton + proton \(\rightarrow\) helium-2 fusion. Stars would rapidly exhaust their fuel and collapse to degenerate states or black holes. In contrast, if \(\alpha_s\) were decreased by about 10% then the deuterium nucleus would cease to be bound and the nuclear astrophysical pathways to the build up of biological elements would be blocked. Again, the conclusion is that there is a rather small region of parameter space in which the basic building blocks of chemical complexity can exist.

We should stress that conclusions regarding the fragility of living systems with respect to variations in the values of the constants of Nature are not fully rigorous in all cases. The values of the constants are simply assumed to take different constant values to those that they are observed to take and the consequences of changing them one at a time are examined. However, if the different constants are fully linked together, as we might expect for many of them if a unified Theory of Everything exists, then many of these independent variations may not be possible. The consequences of a small change in one constant would have further necessary ramifications for the allowed values of other constants. One would expect the overall effect to
be more constraining on the allowed variations that are life-supporting. For examples of such coupled variations in string theories see refs. [29, 30, 31].

These considerations are likely to have a bearing on interpreting any future quantum cosmological theory. Such a theory, by its quantum nature, will make probabilistic predictions. It will predict that it is 'most probable' that we find the universe (or its forces and constants) to take particular values. This presents an interpretational problem because it is not clear that we should expect the most probable values to be the ones that we observe. Since only a narrow range of the allowed values for, say, the fine structure constant will permit observers to exist in the Universe, we must find ourselves in the narrow range of possibilities which permit them, no matter how improbable they may be [32], [19]. This means that in order to fully test the predictions of future Theories of Everything we must have a thorough understanding of all the ways in which the possible existence of observers is constrained by variations in the structure of the universe, in the values of the constants that define its properties, and in the number of dimensions it possesses.

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