

Future climates: Markov blankets and active inference in the biosphere

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We formalise the Gaia hypothesis about the Earth climate system using advances in theoretical biology based on the minimization of variational free energy. This amounts to the claim that nonequilibrium steady-state dynamics—that underwrite our climate—depend on the Earth system possessing a Markov blanket. Our formalization rests on how the metabolic rates of the biosphere (understood as Markov blanket’s internal states) change with respect to solar radiation at the Earth’s surface (i.e., external states), through the changes in greenhouse and albedo effects (i.e., active states) and ocean-driven global temperature changes (i.e., sensory states). Describing the interaction between the metabolic rates and solar radiation as climatic states—in a Markov blanket—amounts to describing the dynamics of the internal states as actively inferring external states. This underwrites climatic nonequilibrium steady-state through free energy minimisation and thus a form of planetary autopoiesis.

Key words: *Autopoiesis, active inference, free energy minimization, Earth’s climate system, Gaia hypothesis*

1- Introduction

The standard models of the Sun’s evolution show an increase in its radiation and brightness over time [1,2]. Yet, there is empirical and model evidence that, despite exposure to increasing radiation, the temperature of Earth’s climate has remained bounded at habitable levels $\approx 0\text{--}40^\circ\text{C}$ since the Archean (4 billion years before present) [3,4]. In contrast, Earth’s climate dynamics would not be possible for its planetary Lifeless neighbours—such as Mars and Venus—whose dynamics are non-habitable [5].

It has been noted since the inception of physiology that biological systems maintain their organization and bounded internal conditions in the face of external fluctuations [6]. In contrast, the entropy of inert (and closed) systems is unbounded and increases indefinitely. This asymmetry between external and internal conditions is thus broadly recognized as an important characteristic of biological systems [7,8]. Schrödinger asked about such internal conditions, ‘*how can the events in space and time which take place within the spatial boundary of a living organism be accounted for by physics and chemistry?*’ [9], p.2.

Recent advances in theoretical biology suggest that biological systems can resist dissipation and external fluctuations through predictive behaviour [10–12]. That is, biological systems preserve their organization and bounded internal conditions by anticipation or active inference, or at least behave as if they had these predictive faculties. The underlying observation is that the time evolution of all systems, whether they are biological or not, depends on the past. However, the time evolution of living systems, looks as if it depends not only on the past and present, but also on the future [10–12]. The reason for this is that living systems act on the basis of a predictive model of their ambiance: they appear to model their ambiance to preserve their organization and bounded internal conditions [10–12]. Such predictive behaviour is named anticipation [10,13–16], allostasis [11,17,18] and recently, active inference [12,19–21]. Active inference is a corollary of the free energy principle that describes the self-organisation of systems that can be distinguished from their external milieu, in virtue of possessing a Markov blanket [12,21].

Since the asymmetry between the Earth’s internal and external conditions implies some sort of action to maintain the bounded temperatures at habitable levels, we therefore offer the following hypothesis: can the Earth’s climate system be

47 interpreted as an anticipatory system that minimizes variational free energy? [22]. This hypothesis provides an interesting
48 connection with the Gaia hypothesis which argues for Earth's planetary bounded internal conditions *by* and *for* the biosphere
49 [23–25]. Here, we formalise the Gaia hypothesis by providing key organizational relationships among the atmosphere,
50 hydrosphere, lithosphere, and biosphere that allow a mathematical formulation of a Markov blanket for the Earth's climate
51 system. This should be considered as a precondition for interpreting and proving (elsewhere) that the temperature of the
52 Earth's climate—bounded within habitable ranges—has arisen due to an anticipatory or active (Bayesian) inference. In the
53 following section, we briefly review the relationship between Markov blankets, free-energy minimization, and active inference.
54 We then propose, based on empirical evidence and prior model-based theoretical work, the existence of a Markov blanket for
55 the Earth's climate system. Finally, we point out some implications of this formal treatment for studying the Earth's climate
56 dynamics.

57 **2- Free energy, Markov blankets and active inference**

58

59 Although the principle of free energy minimization arose in neuroscience, it turned out to be sufficiently generic to
60 ascribe cognitive processes to all living systems [12]. Thus, the minimization of free energy (a.k.a., a generalised prediction
61 error) can be regarded as a dynamical formalization of the embodied cognition implicit in autopoiesis or biological self-
62 production [12,26–30]. One of the conditions for the existence of an autopoietic system is a boundary [31], which resonates
63 with Schrodinger's question above. A Markov blanket defines the boundary between a system of interest and its environment in
64 a statistical sense. More specifically, it provides a statistical partitioning of a system into states internal and states external to
65 the system. In this context, a Markov blanket is a set of variables through which things internal and external to a system
66 interact.

67

68 Important examples of Markov blankets arise in multiple fields. One of the most commonly encountered is the
69 present – which is the Markov blanket separating the past from the future and underwrites the notion of a Markov process.
70 Markov processes (Gagnic 2017) are stochastic (random) processes whose dynamics may be characterized without reference
71 to their distant past. In other words, if we know the present state of a system, knowing about the past tells us nothing new about
72 the likely future. On one reading of Newtonian physics, the positions and momenta of particles can be seen as the Markov
73 blankets through which particles interact with one another (Friston 2019). In the life sciences, Markov blankets have been
74 associated with the physical boundaries surrounding cells (Friston 2013, Kirchhoff, Parr et al. 2018, Palacios, Razi et al. 2020)
75 (i.e., their membrane) through which all influence between the intracellular and extracellular spaces are mediated. At larger
76 spatial scales, they have been drawn around plant physio-anatomy (Calvo and Friston 1917) and neural networks (Hipolito,
77 Ramstead et al. 2020) such that the superficial and deep pyramidal cells of the cerebral cortex play the role of a Markov
78 blanket, through mediating interactions between different cortical columns. This concept has been extended to an arbitrary
79 number of spatial scales (Ramstead, Badcock et al. 2018), with muscles and sensory receptors acting as an organism's
80 Markov blanket, and specific organisms acting as blankets to separate groups of organisms. The thing all of these examples
81 have in common is that they separate the world into two sets of states, which interact only via their Markov blanket (Clark
82 2017).

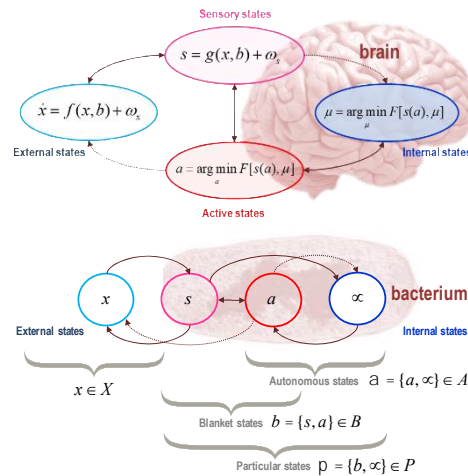
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84 Pearl [32] introduced the term 'Markov blanket' to describe the set of variables that mediate relationships to and from
85 a variable of interest. More precisely, for a given variable X and its known blanket states b , no more information about X is
86 gained when knowing the state of variables outside b . This does not imply that knowing the blanket states allows one to fully
87 predict the evolution of the target variable. This is because there may be some intrinsic properties in the system that renders X
88 stochastic. Markov blankets segregate 'directed graphs' known as Bayesian networks such that one side of the blanket is
89 conditionally independent of the other, given the blanket. This conditional independence clearly has important existential
90 implications.

91

92 The Markov blankets that one considers in active inference, comprise sensory and active states, that feature specific
93 coupling or causality relationships with internal and external states [12,21] (see Figure 1). Here, the term "states" stands for any
94 variable of the system. A stipulative condition for the existence of a Markov blanket is that internal states must be conditionally

95 independent of external states, given blanket states and vice-versa. This formalizes the idea that there are no direct
 96 interactions between internal and external states, only via the blanket states. In other words, internal and external states can
 97 only influence one another via sensory and active states and the internal states only ‘see’ the external states through the ‘veil’
 98 of the Markov blanket [12,21]. Specifically, active states mediate the influence of internal states on external states, and sensory
 99 states mediate the reciprocal influences. In other words, internal states cannot influence sensory states, while external states
 100 cannot influence active states [12,21] (Figure 1). The conditional independence between internal and external states—provided
 101 by blanket states—enables interactions between the inside and outside of the system, but only via the blanket. It is worth noting
 102 that the terms ‘active’ and ‘sensory’ derive from applications of this formalism to biology. However, the same mathematical
 103 relations exist in Hamiltonian dynamics, where these may be replaced with the labels ‘position’ and ‘momentum’.



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 105 **Figure 1. Markov blankets and active inference.** Panel A and B illustrates the partition of states into internal states and hidden or
 106 external states that are separated by a Markov blanket—comprising sensory and active states in the brain and in a cell, respectively. For the
 107 assignment of states in the brain see reference [33,34]. B) The internal states can be associated with the intracellular states of a cell, while
 108 sensory states become the surface states of the cell membrane overlying active states (e.g., the actin-like MreB filaments [REF] that mediate
 109 cell mobility). The intracellular state dynamics of a cell correspond to perception, while action mediates coupling from internal states to
 110 external states. Importantly, once such a Markov blanket is established for a system of interest, along with the necessary condition of non-
 111 equilibrium steady state (NESS), one can formally show that the internal states predict external states and thereby autoregulate the system,
 112 via a minimization of variational free energy [33,34]. This in turn implies that the expected entropy of sensory states remains bounded,
 113 thereby ensuring resistance to dissipation by external fluctuations. Note that s here is assumed to be given by a static function of x . This
 114 contrasts with the dynamical formulation presented in some papers. The underlying reason for this is that most software implementations
 115 use g to specify a likelihood of sensations given external states, under an adiabatic approximation to the underlying dynamics.

116 Active inference is, thus, an account of autopoiesis in dynamic terms, provided that such systems are (i) at non-
 117 equilibrium steady state (NESS) and (ii) that can be statistically segregated from their environment as in Figure 1. The first
 118 NESS condition simply means that a system persists over a nontrivial time-scale and does not dissipate. This implies the
 119 existence of a NESS density to which the system self-organises that can be thought of as a probabilistic description of the
 120 system’s pullback attractor [35,36]. The second (statistical segregation) condition implies the presence of a Markov blanket
 121 [20,32]. The NESS dynamics of the system—and the presence of the conditional independencies implied by a Markov
 122 blanket—means that the average internal state effectively parameterise a probability distribution over the external states
 123 [12,20,33,34]. In other words, for any given blanket state, the average internal state represents the causes (i.e., hidden, or
 124 external) of sensory impressions. It is fairly straightforward to show that the dynamics of the (average) internal state constitute a
 125 gradient flow on a variational free energy functional of this parameterised density. The minimization of free energy (and implicit
 126 minimisation of sensory entropy) enables biological systems to maintain their sensory states within physiological bounds, and
 127 undertake predictive behaviour about the causes of their sensation necessary to sustain their existence [12,21,37].

128
 129 In this context, minimisation of free energy means resisting entropic fluctuation and maintenance of a bounded set of
 130 physiological states. As free energy is a functional of a probabilistic model (the parameterised density over external states

131 above), this means a system must instantiate an implicit model of its ambient space. Minimising free energy fits such a model
132 to sensory states, thereby ensuring good predictive behaviour. In Bayesian statistics, the evidence for such a model is known
133 as the ‘model’ evidence or marginal likelihood: namely, the probability of observing some data, given a model of how those data
134 were generated. Variational free energy upper bounds the negative log model evidence, which is a ubiquitous quantity in
135 statistical physics, Bayesian statistics and machine learning [38]. In machine learning, the variational free energy is commonly
136 called the evidence lower bound, or ELBO [39]. In engineering, it is the cost function associated with Kalman filters. In
137 information theory, minimising free energy corresponds to maximising efficiency or minimum description length approaches. In
138 predictive coding, the evidence is taken as the (precision weighted) prediction error. Crucially, in the free energy framework
139 these are all the same thing: the probability distribution encoded by the internal states that quantifies the dynamics of the
140 external states and evolves towards the variational free energy minimum [40]—an upper bound on surprise or negative log
141 evidence (i.e., the negative log probability of finding the system in a particular state) [12].
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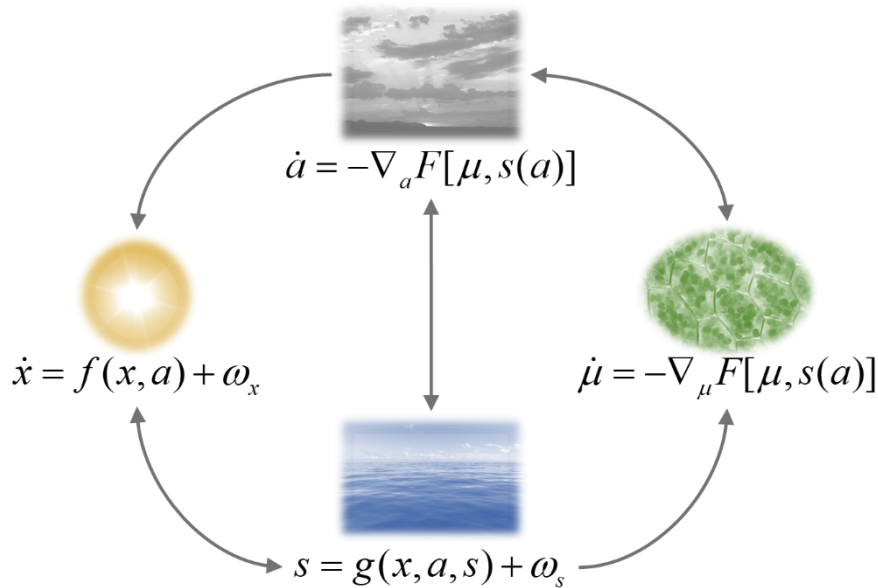
143 The surprise is a function of the states of the Markov blanket itself [40]. Free energy is a function of probabilistic
144 beliefs, encoded by internal states about external states (i.e., expectations about the probable causes of sensory interaction),
145 given any blanket state¹. When these beliefs are equal to the posterior probability over external states, free energy becomes
146 equivalent to surprise. Otherwise, it is always greater than (i.e., constitutes an upper bound on) surprise [40]. Vicarious
147 interaction through a Markov blanket lends an interpretation to the dynamics of such systems, as if internal states were inferring
148 external states based upon the blanket states. This implies that the kind of organization of Markov blankets we consider results
149 in processes that work to seek out evidence (active inference), namely self-evidencing dynamics underlying the autopoietic—
150 thus autonomous—organization of life.

151 In brief, this means we can characterize living systems as minimising variational free energy, and therefore surprise,
152 where the minimisation of variational free energy entails the optimisation of beliefs about things beyond the Markov blanket
153 (i.e., external states). Thus, external fluctuations can therefore (or must therefore) be modelled to maintain a bounded set of
154 physiological states. This formalism has been applied to a range of problems in biology [12,28,41,42] including thermodynamic
155 physical systems [33,34]. On the basis of the ensuing Bayesian process of active inference, we turn next to defining and
156 evidence-based Markov blanket for the Earth’s climate system.

157 **3. Markov blankets and the Earth’s climate system**

158 Several authors suggest that autopoiesis happens at the planetary scale [43–52]. The implication of this view is that
159 minimization of variational free energy must also occur at the planetary level. The challenge for this perspective is to identify the
160 Markov blanket of a climate system (here, the Earth) such that we can think about how internal states of the system may
161 appear to infer and act upon external states. Delineating this blanket is essential in finding the generative model that
162 determines climate dynamics. To do so, we need to define variables (i.e., states) internal and external to the Earth’s climate
163 system—those states that constitute or are intrinsic to the system and those that are not. In so doing, we regard empirical and
164 model-evidence based interactions between the Earth’s climate components and identify a set of variables that satisfy the
165 conditional independencies allowed by the Markov blankets that underwrite active inference (Figure 2):
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¹ Technically, this means that surprise is a *function* of blanket states, while variational free energy is a *functional* of a probability distribution about external states, given a blanket state.



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Figure 2 – Earth’s climate and Markov blankets. This figure illustrates the conditional dependencies between four key variables in the Earth’s climate system. These conditional dependencies imply a Markov blanket comprising active (a) and sensory (s) states (in analogy with perception-action loops in cognitive science) that mediate the interactions between internal (μ) and external (x) states. The internal states are the metabolic rate changes of atmospheric greenhouses and aerosols turnover (see section 3.2 for the full explanation). These influence active states (changes in greenhouse and albedo effects) (see section 3.3) and vicariously the external states (solar radiation changes at the Earth’s surface) (see section 3.1). The external states cause changes in the sensory states (ocean-driven global temperature changes) (see section 3.4), which leads to changes in the internal states. External (and sensory) states exhibit dynamics prescribed by stochastic differential equations (or a static stochastic mapping) with Gaussian white noise uncorrelated in time (ω). Internal and active states perform a gradient descent on variational free energy (F), defined in relation to a (characteristic) NESS density for the external and sensory states. As a consequence, the internal states appear to model and act upon the external and sensory states.

Figure 2 illustrates the conditional dependencies between variables forming the Earth’s climate system and Markov blanket, highlighting that the metabolic rates of the biosphere (internal states) only influences solar radiation at the Earth’s surface (external states) via changes in greenhouses and albedo effects (active states), while reciprocal interactions are mediated by the ocean-driven global temperature changes (sensory states). We turn next to defining each of these variables that constitute this climatic partition of states.

3.1 External states (x): Solar radiation changes at the Earth’s surface

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The space weather² and external environment of the Earth system, such as the electromagnetic solar radiation and galactic cosmic shortwave radiation and energetic electron precipitation, are the source of energy, but also of external forcing and perturbation, respectively [53]. In many mainstream climate models (radiative energy balance models, EMICs and GCMs) cosmic rays are not a source of forcing fluctuation in the climate system. Indeed, those who normally study the impact of cosmic rays on Earth are not climatologists, but astrophysicists and planetary scientists. In line with the interdisciplinary perspective offered here, we mention these possibilities – which could be incorporated into the external states of a Markov blanket for the Earth’s climate system. However, the key construct of the Markov blanket does not depend upon this, and the external states stand in for anything outside of the system of interest that causes changes to the blanket states.

² “Space weather refers to dynamic conditions on the Sun and in the space environment of the Earth, which are often driven by solar eruptions and their subsequent interplanetary disturbances” [56].

197 According to the standard model of stellar evolution, the incoming solar radiation during the Archean to the
198 Proterozoic eon (3.8 billion to 542 million years ago) was 20–30% lower than at present [1,54]. The faint young Sun was only
199 76%-83% as intense as its current value [55]. Also, the Sun often displays extreme and severe coronal mass ejections, solar
200 flares, and storms, making the Earth's space weather difficult to predict [56]. Sunspots and low solar activity (solar Maunder
201 minimum) correspond with historically documented cold periods on Earth—and is often related to little ice ages, in the form of
202 frost fairs [57]. Diverse reconstructions of past climate records, in cosmogenic isotope archives, have revealed associations
203 between the Earth's climatic response to solar radiation changes and cosmic ray variations [58].
204

205 Despite changing space weather, such as solar radiation fluctuation, the Earth is close to being in *radiative*
206 *equilibrium*. The incoming energy (mainly of the Sun) is balanced by an equal flow of heat that the Earth radiates back to outer
207 space. Under the condition of the Earth's so-called energy flux balance [59], Earth's temperature is bounded at habitable levels
208 [4]. How the energy flows into—and away from—the Earth is key to understanding climate dynamics [60]. This also indicates
209 that although the solar radiation at the Earth's surface is necessary, it is not the only factor affecting global temperatures on
210 Earth. There are different constraints and processes that account for the Earth's energy flux balance. These depend on the time
211 scale. At very short times scales, among other phenomena, the incoming Sun radiation leads to evaporation of seawater and
212 sea surface temperature changes [61]. At millennial times scales (the last 3 million years) changes of solar radiation at the
213 Earth's surface due to variations in the Earth's orbit *triggered* climatic changes such as glacial-interglacial oscillations
214 (variations in ice volume and ice sheet extent) [62], the so-called Milankovic effect of orbital (astronomical) forcing. At long time
215 scales (from 2 to 4 billion years), different constraints on the Earth system offset the weak forcing and radiation of the faint
216 young Sun. Thus, the solar radiation exerts different forcing at different time-scales. Our proposal is that changes in solar
217 radiation at the Earth's surface account for the external states of the Earth's Markov blanket (x) (Fig. 2).
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219 For the Earth's Markov blanket proposed here, volcanoes—a source of carbon and abiogenic aerosols—may not be
220 external to the Earth's climate system. This follows from the next argument: the structure of inner and outer layers of the
221 continental plates appears to have been dominated by water-dependent continental drift changes [63]. The hydrological cycle,
222 which allows continental drift and geomorphological changes, is life-dependent [64]. Together with the shift from the reductive
223 to oxidative atmosphere—from the advent of photosynthesis and the shifting balance between seafloor and terrestrial
224 weathering REFMills et al in the middle Archean to the proterozoic period —resulted in the strength of the hydrological cycle,
225 continental drift, continent growth, plate spreading, volcanic activity [65–68]. Tectonic and volcanic activity and their products,
226 therefore, are ultimately internal to the Earth system functioning, and thus to its Markov blanket.
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228 Under free energy minimization, the forcing by the external sates may affect Earth's climate only through its Markov
229 blanket's sensory states ($x \rightarrow s$) (ocean-driven global temperature changes) (Fig. 2). At first glance, one might think this to be
230 only partially true, because the solar irradiance impacts on the Earth's upper atmosphere. However, the insolation on the upper
231 atmosphere is quite different to the insolation reaching the Earth's surface after passing through the atmosphere, most of which
232 is metabolically derived (see next sections). Hence, changes of atmospheric states triggered by the solar irradiance and cosmic
233 ray particle ionization—such as large-scale electric current dynamic changes in the troposphere [69], or enhanced cloud
234 formation [70]—are ultimately determined by the internal states through active states. In short, the changes in solar radiation at
235 the Earth's surface may be selected by the internal states through active inference.

236 **3.2 Internal states (μ): metabolic rate changes**

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238 Internal states must be conditionally independent of external states provided blanket states and vice-versa. This
239 formalises the idea that there are no direct interactions between internal and external states, only via the blanket states. Here
240 we propose that the metabolic rate changes fulfil this condition, and thus are the Earth's Markov blanket internal states (μ). We
241 base this proposition on the following arguments.
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243 The metabolic rate reactions directly depend on habitable bounded temperatures and only indirectly on the incoming
244 solar radiation (external states). This means that the external states of Earth's Markov blanket can only affect the metabolic rate
245 changes through bounded sensory states ($x \rightarrow s \rightarrow \mu$) (Fig 2). One may argue that this is not true, because the photosynthesis

246 depends on solar radiation. However, at low temperatures, between 0-10°C or above 20°C the enzymes that carry out
247 photosynthesis do not catalyse at the optimum photosynthetic rate, which is obtained between 10 and 20°C [71,72]. Indeed, all
248 the metabolic rate reactions on the Earth system depend directly on the physiological temperatures between ≈0–40°C [74] at
249 which the Earth has remained since the Archean [3].

250 The metabolic rate changes affect the incoming solar radiation (external states) only through its active states ($\mu \rightarrow$
251 $\alpha \rightarrow x$) (Fig 2), which are fundamentally linked to the rate of net primary productivity (NPP). While the biosphere captures no
252 more than 1% of solar radiation, and thus can be considered insignificant on the Earth's energy flux balance, from the Archean
253 to the present the atmospheric and lithospheric chemical elements have been continuously metabolically transformed,
254 produced [25,75–79], mobilized [80], localized [81], and integrated into biogeochemical cycles by a relatively stable microbial
255 set of core enzymes involved in major redox reactions and electron transfer of Earth's chemistry, despite enormous genetic
256 diversity [82]. Indeed, through Earth's ontogeny global NPP had contributed three times more energy to the geochemical cycles
257 than Earth's internal heat [65]. While the contribution of the marine ecosystems (mostly microbial) is longer over geological time
258 scales, the continental ecosystems per unit area contributes much more and faster due to availability of key nutrients and solar
259 energy. However, due to the extension of the ocean and the accelerated rates of continental weathering in the continents both
260 have equivalent contribution to the global NPP. Thus, the metabolic rate changes have resulted, among other things, in
261 changes in greenhouse and albedo effects (active states) (Fig. 2), and hence in the Earth's energy flux balance and habitable
262 temperatures.

263 More specifically, the biogeochemical carbon cycle, upon which most of the current climate change depends (IPCC
264 2013), involves carbon dioxide (CO₂) and methane (CH₄) in the troposphere due respectively to metabolic enhancement of rock
265 weathering [83,84] and methanogenesis rates [85]. On Earth 99.9% of CH₄ is due to methanogenesis [86]. The biogeochemical
266 carbon cycle also is linked to the O₂ levels (availability of free energy) and the formation of ozone (O₃) in the stratosphere which
267 intercepts, absorbs and converts more than 97% of the sun's mutagenic ultraviolet radiation into heat [67]. The biogeochemical
268 sulphur cycle involves the production and release into the troposphere of biogenic cloud-forming aerosols, such as dimethyl
269 sulphide (DMS). On average, the aerosols of ocean microorganisms boost the number of cloud droplets by about 60% annually
270 [87]. DMS is produced not just on the sea [88], but also on continents [89,90]. The microbes suspended in the atmosphere also
271 induce cloud formation [91,92]. A biogenic aerosol-climate linkage has been postulated, through the albedo effect of clouds
272 affecting the incoming solar radiation, hence the Earth's energy flux balance and temperature [93]. Model-based evidence
273 shows this is highly plausible [94–96]. What is crucial is that the existence of glaciers and ice sheets, which translates
274 effectively into changes in albedo effects (active states), depends on the accumulation of snow and therefore on a great part of
275 global cloud cover derived from metabolism.

276 Metabolism is a self-organizing phenomenon [97–101] involving not just the biosphere, but the atmosphere,
277 hydrosphere, and geosphere [103–106]. Nevertheless, the internal states proposed here focuses on carbon cycling, biogenic
278 aerosols and (oxygenic) photosynthesis, whereas the metabolic network of the biosphere is much more complicated [82] and
279 rates may be limited by other components such as nitrogen availability (also biologically constrained). A more extensive
280 overview of the different ways in which metabolic interactions within the biosphere intertwined with the dynamics of the Earth's
281 geochemistry [82] should be considered in the future for a more detailed account of internal states. Here, we simply
282 acknowledge that the system cannot respond with infinite capacity to appropriate temperature ranges when it also depends on
283 biological cycling of essential nutrients. In this regard the dynamics of internal states' response may be critical.

284
285 Different explanations have associated biological evolution at the "species" level with the metabolic responses [Ref
286 1,2]. These proposals suggested that the persistence of the biosphere increased its likelihood of acquiring further persistence-
287 enhancing traits and that species that destabilize their environment are short-lived and result in extinctions until a stable state is
288 found. While it is reasonable to say that metabolic rate changes plays out on the diversity, richness, abundance, and
289 connectivity (trophic and symbiotic relations) of eukaryotic systems and prokaryotic microorganisms in the ocean, continents,
290 deep subsurface and atmosphere [45,108–110], "speciation" requires the arising of new lineages, which may take ecological
291 and geological timescales. At least, so far, there is no experimental evidence of speciation under controlled conditions even at
292 the level of operational taxonomic units in microorganisms. Thus, although the biospheric diversity encodes the metabolic
293 capacity, it is more congenial that the dynamics of the internal states' responses are associated with metabolic rate changes

294 based on a relatively stable phylogenetic redundancy of a set of a key core enzymes (e.g. Rubisco) [82] [107] scattered in a
295 single rhizome network of the biosphere [102]. Enzymes are fast acting and demand low energy of activation [Ref], which
296 continuously and directly ensures the changes in greenhouse and albedo effects that constitute the active states (Fig. 2), and
297 thus instantiate the internal states (μ) beneath the Earth's Markov blanket.
298

299 **3.3 Active states (a): changes in greenhouse and albedo effects**

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301 From the Archean to the Proterozoic eons (from 4 to 2.5 billion years ago), solar radiation alone was too weak [1,2],
302 to maintain liquid water and ice-free conditions on Earth [54] (see section 3.1). Indeed, if all other global parameters are held
303 constant, and if the Archean-Proterozoic Earth's greenhouse gas concentration in the atmosphere were the same as now (405
304 ppm/0.0405% and 722ppb/0.007% and of the air of CO₂ and CH₄ respectively), the Earth's mean temperature would have been
305 below the freezing point of seawater (−18 °C) [55,111]. Earth would have been in continual deep freeze, with glaciers reaching
306 the Equator, at until one billion years ago, when the solar radiation had increased enough to melt the ice [55,111]. In this case,
307 the internal states of the Earth's system, which rely on liquid water and physiological temperatures, would have been very
308 limited or perhaps non-existent. And yet, there is ample geologic, paleontological and model-based evidence regarding this
309 geological period showing the existence of extensive bodies of liquid water at the surface and continuous habitable Earth's
310 temperatures (≈0–40°C) [3,54,55,112,113]. This puzzle is called the faint young Sun paradox.
311

312 Earth's climate responses compensated for the young Sun's lower radiation at the Archean Earth's surface through
313 the warming produced by higher greenhouse gas concentrations and reduced albedo [3,54,55,112–114]. The so-called
314 greenhouse effect, on which greenhouses gases trap heat emitted from the surface's reflection. Precisely, the CH₄, which has
315 stronger radiative forcing than CO₂ and is almost entirely metabolically produced (see section 3.2), could have provided 10–12
316 °C of surface warming [115] with levels ranging 10² to 10⁴ times higher than modern amounts [3]. A high amount of atmospheric
317 nitrogen (increasing atmospheric pressure), dependent on the biogeochemical nitrogen cycle, would have given 4.5°C extra
318 warming [116]. Thus, the increment of greenhouse gas fluxes and abundance was *necessary* to offset a fainter Sun [3,112], if
319 the habitable conditions of the Earth's system were to be maintained.

320 The solar radiation has steadily increased by twenty to thirty percent from the Mesoarchean, to the Proterozoic, to the
321 present [1,2]. That is, the faint young Sun became a bright mature Sun. Yet, the Earth's liquid water did not evaporate as with
322 Venus or dissipate as with Mars [5]. Increasing solar forcing through time is roughly cancelled by decreasing greenhouse and
323 increasing albedo effects by atmospheric clouds and lithospheric ice-snow cover, which reflect the Sun's radiation back into
324 space [117,118]. Evidence shows that over that period, the CH₄ and CO₂ greenhouse gases declined to their preindustrial
325 values, ~722ppb/0.007% and ~200 ppm/0.02% respectively [114]. This was an active process. Oxidative metabolic enhanced
326 silicate weathering dampened the CO₂ concentration 1000 times faster [83,119] compared to inorganic anoxic carbonate–
327 silicate weathering [120]. The oxidative atmosphere derived from the evolution of photosynthesis shifted the metabolic rates
328 towards oxidation and thus faster reduction of greenhouse gases [68,121]. The Proterozoic CH₄ has been damped by reverse
329 methanogenesis, causing an anti-greenhouse effect by forming a thick hydrocarbon organic haze given the changes of
330 CH₄/CO₂ ratios higher than ~0.1 [115,122]. Moreover, since the late Archean, climate moderation by the carbon cycle (mainly
331 CO₂/CH₄) is consistent with the increase of snow-ice cover, and thus occasional glaciations [3,123]. Glaciers, ice-sheets, and
332 snow cover depend on the cloud formation. Model-based evidence shows that the net effect of cloud cover is cooling [94–96].
333 Glaciers, ice-sheets and snow cover account for more than 70% of the Earth's albedo effect [60]. These results indicate that the
334 albedo effect is necessary to counterbalance the steadily increase of incoming solar radiation.

335 These rather basic observations allow us to assert that changes in greenhouse and albedo effects instantiate the
336 active states of the Earth's Markov blanket. Both effects, the greenhouse and albedo, are conditionally dependent on the
337 metabolic rate changes that result in modelling, inference and selection, through this blanket, of the incoming solar radiation at
338 the Earth's surface (external states) ($\mu \rightarrow a \rightarrow x$) (Fig. 2). Importantly, by acting upon the external states, the active states of

339 the Earth's Markov blanket also bind (i.e., upper bound) the entropy of its sensory states, so that it does not increase
340 indefinitely, i.e., dissipate ($a \rightarrow x \rightarrow s$) (Fig. 2)³.
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342 **3.4 Sensory states (s): ocean-driven global temperature changes**

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344 The ocean is a driving force affecting the Earth's mean temperature and thus the climate system's temporal evolution
345 [60,124,125]. Changes in Earth's mean temperature depend sensitively on the thermal inertia of the ocean's capacity to absorb
346 heat (ocean heat uptake) [126,127]. The ability to absorb heat from the ocean is influenced by the physical-chemical properties
347 of the water (high specific heat capacity), and also by the dynamics of the ocean itself and its interaction with the atmosphere.
348 That is, the ocean does not change its temperature "rapidly" as the atmosphere does, and the oceans' heat uptake reduces the
349 effective climate sensitivity⁴ and weakens its warming response [124–126,128,129]. As a consequence, the ocean drives the
350 global temperatures by distributing heat around the planet, responding very slow to thermal fluctuation and absorbing 1000
351 times more heat than the atmosphere without changing its temperature. It thus accumulates about 93% of the Earth's thermal
352 energy [130]. The excess energy that the ocean stores can lead to the ocean's thermal expansion, e.g., melting of ice sheets
353 and thus to sea-level rise, which on the latest glacial-interglacial oscillation represents changes in sea level over 100 m [131].
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355 These empirical and model-based observations let us consider the ocean-driven global temperature changes as
356 accounting for the sensory states (s) of the Earth's Markov blanket (Fig. 2), and, therefore, bidirectional interactions of the
357 sensory states with external and active states. Incoming solar radiation (external states) mainly causes evaporation of ocean-
358 surface water via sea-surface heat and surface temperature changes [61]. This affects water-vapor concentration in the
359 troposphere, air-temperature distribution, winds, cloud formation and intermittent precipitation on continents and the absorption
360 of greenhouse gases⁵ [132]. All these evince the influence of external on sensory and of sensory states on active states.
361 Conversely, the effect of active states on sensory states is most likely the freshwater nutrient-rich fluxes from continents and
362 ice-sheets to the oceans that create density gradients of salinity [133]. It is through the thermal and density gradients of
363 seawater that sensory states affect (external states) incoming solar radiation changes at the Earth's surface. Both gradients
364 ultimately determine ocean convection, turnover, and general circulation accounting for the slowdown in surface warming and
365 the cooling of the ocean with the relevant effects on the glacial cycles [REF]. The overall ocean dynamics results in moving
366 heat from Earth's equator to the poles, distributing energy throughout the Earth's climate components, and thus on the ocean-
367 driven global temperature changes [124–126,128,129].
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369 These arguments speak to the dynamics of Markov blankets, in which external states cannot directly affect changes
370 in active states (greenhouse and albedo effects), but only as mediated through the sensory states. Similarly, the changes in
371 solar radiation at the Earth's surface affect the metabolic rate changes (internal states) only through the sensory states (ocean-
372 driven global temperature changes) ($x \rightarrow s \rightarrow \mu$) [132,134]. That is, the ocean dynamics becomes the sensory states for the
373 dynamics of whole biosphere (marine and continental) (see section 3.2). At first glance, one might question this, because the
374 oceans cannot mediate the interactions between solar radiation and the continental ecosystems. However, the metabolic rate
375 changes, not just in marine, but in continental ecosystems are driven by the Earth's mean temperature [79,135,136], which in
376 turn is buffered, hence driven by the oceans' heat uptake and dynamics. At the same time, the only way that the internal states
377 affect the ocean is through the chemical modification of the atmosphere and lithosphere, which results in changes of
378 greenhouse and albedo effects (active states) (see previous sections). This interaction between internal and external states as
379 climatic states—in a Markov blanket—in turn will look present as the instantiation of active inference at a planetary scale.
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³ Technically speaking, the entropy of the sensory state is effectively bounded from above and below. This follows, because the active maintenance of an Earth-like nonequilibrium steady-state precludes both high and low entropy NESS densities, e.g., solar, and lunar climates, respectively.

⁴ Climate sensitivity refers to how much, in the near and the long-term, the Earth's climate will warm (or cool) after a perturbation like an increase in CO₂ concentrations or solar radiation.

⁵ The ocean is the most important sink of atmospheric CO₂ [132].

381 **4. Discussion: Active inference at the planetary-scale**

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Within the free energy framework, the consideration of planetary autopoiesis as active inference rests upon the existence of a Markov blanket for the Earth's climate system. That is, in the relation and coupling of internal and external states through the Markov blanket. If we take the perspective that the metabolic rates change (the internal states of the Earth's Markov blanket) is performing active inference about the changes in incoming solar radiation (external states), we can frame their dynamics in terms of approximate Bayesian inference, which is equivalent to the minimization of variational free energy. This treats the metabolic rate changes as if they parametrize an implicit (variational) probability density over solar radiation changes at the Earth's surface and optimize this to maximize model evidence of changes in external states to preserve habitable conditions.

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As the metabolic rate changes influence changes in greenhouse and albedo effects, the indirect or mediated influence of solar-radiation variation at the Earth's surface is mathematically equivalent to Bayesian inference. This is because we could, in principle, use the average solar radiation changes at the Earth's surface to draw inferences about the variables affecting the metabolic rate changes. This perspective implies a NESS density associated with the blanket and its external states, i.e., the density that the system tends toward when it is perturbed. In analogy with biological-like behaviours at the planetary-scale proposed by the Gaia hypothesis, this may be thought of as the range of values within which Earth's homeorhetic dynamics can anticipate (and accommodate) any deviations. Dynamic anticipation of this sort implies that changes in greenhouse and albedo effects also optimize the same quantity through changing ocean-driven global temperatures (directly and via external states).

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An alternative interpretation of the NESS density is as a generative model. This implies that the metabolic rate changes implicitly may model solar radiation changes at the Earth's surface and select them via changes in greenhouse and albedo effects. This process of selection through modelling the environment means that to find the dynamics of the metabolic rate changes—and their influence over changes in greenhouse and albedo effects—we need only to specify the NESS for solar radiation variation at the Earth's surface and ocean-driven global temperatures. The dynamics of active and internal states will emerge from minimizing free energy. That is to say, a sort of planetary selection of permissible perturbations from solar radiation changes at the Earth's surface results in the continuous operating of the Earth's system habitability. What is selected will depend on the active inference at planetary-scale, in relation to the model it generates through the integrated dynamics of the biosphere, atmosphere, lithosphere, and hydrosphere.

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This perspective of such a mathematical formalism may be useful in the modelling of the Earth's climate system as a systemic unity in three ways. The first is in inferring parameters of the implicit NESS density (or generative model) that underwrites the Earth's climate dynamics. The second is that alternative models (with alternative NESS densities) may be proposed to formulate alternative hypotheses about the Earth's climate system. A common mathematical framework enables model comparison (i.e., hypothesis testing) to adjudicate between these hypotheses. Thirdly, this may be useful in predictive modelling, that is, in establishing how different interventions may impact the long-term evolution of Earth's climate. A pragmatic consideration here is that if one can write down the NESS density over external (solar) and sensory (oceanic) states—that is characteristic of the Earth—one can simulate the metabolic (i.e., internal states), greenhouse and albedo effects (i.e., active states) using a gradient flow on variational free energy. The key point here is that the requisite gradients are analytically and numerically tractable; enabling the climactic simulations, much along the lines of how active inference is used to model experimental subjects in cognitive science (neuroscience and ethology).

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While this paper sets out a possible conditional dependency structure between environmental variables, consistent with a Markov blanket, this should be viewed as a hypothesis. This is subject to evidence that could support or refute this. For example, if it were demonstrated that solar radiation at the Earth's surface and metabolic rates in the biosphere are not (approximately) conditionally independent of one another, once greenhouse, albedo, and ocean temperatures are taken into account, this would offer strong evidence against our hypothesis.

428 The relatively abstract treatment here – based upon conditional dependencies – means we have not specified a
429 functional form for the equations of motion linking the dynamics of each climate component. This highlights the work yet to be
430 done to move from a theoretical framing of climate dynamics to a useful model that can be used to test empirical hypotheses.
431 The next two phases of this research will be as follows. First, we need to specify the functional form of the external state
432 dynamics, as a function of the active states, and the form of the sensory states as a function of the internal and active states.
433 This sets up an implicit generative model from the perspective of the active and internal states and means we can test the face
434 validity of this formulation through numerical simulation. In addition, we hope to demonstrate construct validity in relation to
435 established climate models. The second phase will be to fit these numerical simulations implemented with standard software
436 routines (e.g. `spm_ADEM.m`⁶) to climate data. In doing so, we hope to develop a tool that lets us evaluate the evidence
437 afforded by alternative hypotheses about the causes of these data. Under active inference, these hypotheses are typically
438 framed in terms of the parameters of prior beliefs implicit in a generative model (Schwartenbeck and Friston 2016). In other
439 words, we can think of internal states as evolving as if they held beliefs about the external states, and can test hypotheses
440 about what these beliefs might be. The utility of this framing is that our interest is in how biotic and abiotic elements of the
441 climate interact, so it is helpful to be able to pose hypotheses about the one in relation to the other. However, the perspective
442 that the internal states of the Earth's Markov blanket perform inferences about—and act on—the incoming energy radiation
443 through the ocean (sensory) and the greenhouse-albedo effects (active) states may tell us something profound about the
444 character and nature of the Earth.

447 **5. Conclusion**

448 This paper outlines key organizational relationships among the Earth's systemic components—comprising the
449 atmosphere, hydrosphere, lithosphere, and biosphere—that allows one to posit the existence of a Markov blanket for the
450 Earth's climate system. The motivation for this proposal follows the hypothesis that the nonequilibrium steady-state dynamics
451 that underwrite our climate history depends on planetary active inference, which rest upon the existence of a Markov blanket
452 that enshrouds the Earth's internal states. This requires an appeal to the mathematics of living (cognitive) systems, such as
453 minimization of variational free energy, where the formalism of interaction with an external environment has been most
454 comprehensively developed. The Earth system's Markov blanket proposed above conforms to some basic model-based and
455 empirical observations about how the metabolic rate (framed as the internal states) interacts with changes in the solar radiation
456 at the Earth's surface (external states) through ocean-driven global temperatures (sensory states) and through the
457 atmospheric-lithospheric greenhouse-albedo effects (active states). Crucially, a Markov blanket equips the Earth's climate
458 system with a Bayesian process that will allow us to see whether the average internal states appear to engage in active
459 inference—to actively maintain a nonequilibrium steady-state. That is to say, establishing a Markov blanket for the Earth's
460 climate system may allow us to treat its dynamics as performing active inference, in the fashion of biological anticipation
461 (Rosen 1985), and as a form of planetary autopoiesis.

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465 **References**

- 466 1. Gough DO. 1981 Solar interior structure and luminosity variations. In *Physics of Solar Variations*, pp. 21–34. Springer.
- 467 2. Bahcall JN, Pinsonneault MH, Basu S. 2001 Solar models: Current epoch and time dependences, neutrinos, and
468 helioseismological properties. *Astrophys. J.* **555**, 990.
- 469 3. Catling DC, Zahnle KJ. 2020 The Archean atmosphere. *Sci. Adv.* **6**, eaax1420.
- 470 4. Lovelock JE, Kump LR. 1994 Failure of climate regulation in a geophysiological model. *Nature* **369**, 732.

⁶ Part of the freely available as Matlab code in the SPM12 academic software: <http://www.fil.ion.ucl.ac.uk/spm/>

- 471 5. Lammer H *et al.* 2018 Origin and evolution of the atmospheres of early Venus, Earth and Mars. *Astron. Astrophys. Rev.* **26**, 2.
- 472 6. Bernard C. 1878 Lectures on the phenomena common to animals and plants.
- 473 7. Maturana HR, Varela FJ. 1991 *Autopoiesis and cognition: The realization of the living*. Springer Science & Business Media.
- 474 8. Rosen R. 1991 *Life itself: a comprehensive inquiry into the nature, origin, and fabrication of life*. Columbia University Press.
- 475 9. Schrödinger E. 1945 *What is Life?: The Physical Aspect of the Living Cell*. London: The University Press.
- 476 10. Rosen R. 1985 *Anticipatory systems: philosophical, mathematical, and methodological foundations*. New York: Pergamon Press.
- 477 11. Sterling P. 2012 Allostasis: a model of predictive regulation. *Physiol. Behav.* **106**, 5–15.
- 478 12. Friston K. 2013 Life as we know it. *J. R. Soc. Interface* **10**, 20130475.
- 479 13. Nadin M. 2010 Anticipation and dynamics: Rosen’s anticipation in the perspective of time. *Int. J. Gen. Syst.*
480 (doi:10.1080/03081070903453685)
- 481 14. Louie AH. 2012 Anticipation in (M,R)-systems. *Int. J. Gen. Syst.* **41**, 5–22. (doi:10.1080/03081079.2011.622088)
- 482 15. Rosen J, Kineman JJ. 2005 Anticipatory systems and time: a new look at Rosennean complexity. *Syst. Res. Behav. Sci.* **22**, 399–
483 412.
- 484 16. Poli R. 2014 Book review and abstracts. *Int. J. Gen. Syst.* (doi:10.1080/03081079.2014.929869)
- 485 17. Sterling P. 1988 Allostasis: a new paradigm to explain arousal pathology. *Handb. life Stress. Cogn. Heal.*
- 486 18. Schulkin J, Sterling P. 2019 Allostasis: A brain-centered, predictive mode of physiological regulation. *Trends Neurosci.* **42**, 740–
487 752.
- 488 19. Ramstead MJD, Badcock PB, Friston KJ. 2018 Answering Schrödinger’s question: A free-energy formulation. *Phys. Life Rev.*
489 (doi:10.1016/j.plrev.2017.09.001)
- 490 20. Kirchhoff M, Parr T, Palacios E, Friston K, Kiverstein J. 2018 The markov blankets of life: Autonomy, active inference and the
491 free energy principle. *J. R. Soc. Interface* (doi:10.1098/rsif.2017.0792)
- 492 21. Karl F. 2012 A free energy principle for biological systems. *Entropy* **14**, 2100–2121.
- 493 22. Rubin S, Crucifix M. 2017 Is the climate system an anticipatory system that minimizes free energy? *EGUGA* , 16173.
- 494 23. Lovelock J. 1979 *Gaia, a new look at life on earth*. London: Oxford University Press.
- 495 24. Lovelock JE, Margulis L. 1974 Atmospheric homeostasis by and for the biosphere: the Gaia hypothesis. *Tellus* **26**, 2–10.
- 496 25. Margulis L, Lovelock JE. 1974 Biological modulation of the Earth’s atmosphere. *Icarus* **21**, 471–489.
- 497 26. Maturana HR, Varela FJ. 1980 *Autopoiesis and cognition: The realization of the living*. Boston: D. Reidel (originally published in
498 Spanish in 1973).
- 499 27. Allen M, Friston KJ. 2018 From cognitivism to autopoiesis: towards a computational framework for the embodied mind. *Synthese*
500 (doi:10.1007/s11229-016-1288-5)
- 501 28. Friston K, Levin M, Sengupta B, Pezzulo G. 2015 Knowing one’s place: a free-energy approach to pattern regulation. *J. R. Soc.*
502 *Interface* **12**, 20141383.
- 503 29. Bateson G. 1979 *Mind and nature: A necessary unity*. Bantam Books New York.
- 504 30. Von Foerster H. 1984 *Observing Systems*. New York: Intersystems Publications.
- 505 31. Maturana HR. 1980 Autopoiesis: reproduction, heredity and evolution. In *Autopoiesis, dissipative structures and spontaneous*
506 *social orders, AAAS Selected Symposium 55 (AAAS National Annual Meeting, Houston TX, 3–8 January 1979)*, pp. 45–79.
507 Westview Press.
- 508 32. Pearl J. 1998 Graphical models for probabilistic and causal reasoning. In *Quantified representation of uncertainty and*
509 *imprecision*, pp. 367–389. Springer.
- 510 33. Friston K. 2019 A free energy principle for a particular physics. *arXiv:1906.10184*
- 511 34. Parr T, Da Costa L, Friston K. 2020 Markov blankets, information geometry and stochastic thermodynamics. *Philos. Trans. R.*
512 *Soc. A* **378**, 20190159.
- 513 35. Crauel H. 1999 Global random attractors are uniquely determined by attracting deterministic compact sets. *Ann. di Mat. pura ed*
514 *Appl.* **176**, 57–72.
- 515 36. Crauel H, Flandoli F. 1994 Attractors for random dynamical systems. *Probab. Theory Relat. Fields* **100**, 365–393.
- 516 37. Da Costa L, Parr T, Sajid N, Veselic S, Neacsu V, Friston K. 2020 Active inference on discrete state-spaces: a synthesis. *arXiv*
517 *Prepr. arXiv2001.07203*
- 518 38. Friston K. 2010 The free-energy principle: a unified brain theory? *Nat. Rev. Neurosci.* **11**, 127–138.
- 519 39. Bishop CM. 2006 *Pattern recognition and machine learning*. springer.
- 520 40. Friston K. 2018 Does predictive coding have a future? *Nat. Neurosci.* **21**, 1019–1021.

- 521 41. Calvo P, Friston K. 2017 Predicting green: really radical (plant) predictive processing. *J. R. Soc. Interface* **14**, 20170096.
- 522 42. Ramstead MJD, Badcock PB, Friston KJ. 2018 Answering Schrödinger's question: A free-energy formulation. *Phys. Life Rev.* , 1–16. (doi:10.1016/j.plrev.2017.09.001)
- 523 43. von Foerster H. 1975 Gaia's Cybernetics Badly Expressed. *Coevol. Q.* **7**, 51.
- 524 44. Jantsch E. 1980 *The self-organizing universe: scientific and human implications of the emerging paradigm of evolution*. Pergamon Press. See <https://books.google.be/books?id=qx8PAQAIAAJ>.
- 525 45. Margulis L, Sagan D. 1986 *Microcosmos: Four Billion Years of Evolution from Our Microbial Ancestors*. Los Angeles: University of California Press. See <https://books.google.be/books?id=qs4PAQAAMAAJ>.
- 526 46. Margulis L, Sagan D. 1995 *What is Life?* Michigan: Simon & Schuster. See <https://books.google.be/books?id=4IIPQAAMAAJ>.
- 527 47. Onori L, Visconti G. 2012 The GAIA theory: from Lovelock to Margulis. From a homeostatic to a cognitive autopoietic worldview. *Rend. Lincei* **23**, 375–386.
- 528 48. Sahtouris E. 1996 The Gaia controversy: a case for the Earth as an evolving system. In *Gaia in action: Science of the Living Earth* (ed P Bunyard), pp. 324–337. New York: Floris Books.
- 529 49. Capra F, Luisi PL. 2014 *The Systems View of Life: A Unifying Vision*. Oxford: Cambridge University Press. See <https://books.google.be/books?id=iEwHAWAAQBAJ>.
- 530 50. Levchenko VF, Kazansky AB, Sabirov MA, Semenova EM. 2012 Early Biosphere: Origin and Evolution. In *The Biosphere*, InTech.
- 531 51. Rubin S, Crucifix M. 2019 More than planetary-scale feedback self-regulation: A Biological-centred approach to the Gaia Hypothesis.
- 532 52. Clarke B. 2020 *Gaian systems: Lynn Margulis, neocybernetics, and the end of the Anthropocene*. Minneapolis: University of Minnesota Press.
- 533 53. Mironova IA *et al.* 2015 Energetic particle influence on the Earth's atmosphere. *Space Sci. Rev.* **194**, 1–96.
- 534 54. Sagan C, Mullen G. 1972 Earth and Mars: Evolution of atmospheres and surface temperatures. *Science (80-.)*. **177**, 52–56.
- 535 55. Goldblatt C, Zahnle KJ. 2011 Faint young Sun paradox remains. *Nature* **474**, E1.
- 536 56. Liu YD *et al.* 2014 Observations of an extreme storm in interplanetary space caused by successive coronal mass ejections. *Nat. Commun.* **5**, 1–8.
- 537 57. Lockwood M, Owens M, Hawkins E, Jones GS, Usoskin I. 2017 Frost fairs, sunspots and the Little Ice Age. *Astron. Geophys.* **58**, 2–17.
- 538 58. Kirkby J. 2007 Cosmic rays and climate. *Surv. Geophys.* **28**, 333–375.
- 539 59. Stephens GL *et al.* 2012 An update on Earth's energy balance in light of the latest global observations. *Nat. Geosci.* **5**, 691–696.
- 540 60. Stocker TF *et al.* 2013 Climate change 2013: The physical science basis. *Contrib. Work. Gr. I to fifth Assess. Rep. Intergov. panel Clim. Chang.* **1535**.
- 541 61. Liang X, Wu L. 2013 Effects of solar penetration on the annual cycle of sea surface temperature in the North Pacific. *J. Geophys. Res. Ocean.* **118**, 2793–2801.
- 542 62. Milankovitch MM. 1941 Canon of insolation and the iceage problem. *K. Serbische Akad. Beogr. Spec. Publ.* **132**.
- 543 63. Lowman PDJ, Lowman PD. 2002 *Exploring space, exploring Earth: New understanding of the Earth from space research*. Boston: Cambridge University Press.
- 544 64. Harding S, Margulis L. 2009 Water Gaia: 3.5 thousand million years of wetness on planet Earth. *Gaia Turmoil Clim. Chang. Biodepletion, Earth Ethics an Age Cris.* , 41–59.
- 545 65. Rosing MT, Bird DK, Sleep NH, Glassley W, Albarede F. 2006 The rise of continents—an essay on the geologic consequences of photosynthesis. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* **232**, 99–113.
- 546 66. Hawkesworth CJ, Kemp AIS. 2006 Evolution of the continental crust. *Nature* **443**, 811–817.
- 547 67. Falkowski PG. 2006 Tracing Oxygen 's Imprint on Earth 's Metabolic Evolution. *Science (80-.)*. **5768**, 1724–1726.
- 548 68. Kump LR. 2008 The rise of atmospheric oxygen. *Nature* **451**, 277–278.
- 549 69. Lam MM, Tinsley BA. 2016 Solar wind-atmospheric electricity-cloud microphysics connections to weather and climate. *J. Atmos. Solar-Terrestrial Phys.* **149**, 277–290.
- 550 70. Marsh ND, Svensmark H. 2000 Low cloud properties influenced by cosmic rays. *Phys. Rev. Lett.* **85**, 5004.
- 551 71. Grimaud GM, Mairet F, Sciandra A, Bernard O. 2017 Modeling the temperature effect on the specific growth rate of phytoplankton: a review. *Rev. Environ. Sci. Bio/Technology* **16**, 625–645.
- 552 72. Schaum C-E *et al.* 2017 Adaptation of phytoplankton to a decade of experimental warming linked to increased photosynthesis.

- 571 *Nat. Ecol. Evol.* **1**, 1–7.
- 572 73. Puente-Sánchez F *et al.* 2018 Viable cyanobacteria in the deep continental subsurface. *Proc. Natl. Acad. Sci.* **115**, 10702–10707.
- 573 74. Gillooly JF, Brown JH, West GB, Savage VM, Charnov EL. 2001 Effects of size and temperature on metabolic rate. *Science* (80-
574). **293**, 2248–2251.
- 575 75. Margulis L, Lovelock JE. 1975 The Atmosphere as circulatory system of the biosphere—the Gaia hypothesis. *Coevol. Q.*
576 5(Summer) , 31–40.
- 577 76. Izon G, Zerkle AL, Williford KH, Farquhar J, Poulton SW, Claire MW. 2017 Biological regulation of atmospheric chemistry en
578 route to planetary oxygenation. *Proc. Natl. Acad. Sci.* **114**, E2571–E2579. (doi:10.1073/pnas.1618798114)
- 579 77. Atekwana EA, Slater LD. 2009 Biogeophysics: A new frontier in Earth science research. *Rev. Geophys.* **47**, 1–30.
580 (doi:10.1029/2009rg000285)
- 581 78. Jelen BI, Giovannelli D, Falkowski PG. 2016 The role of microbial electron transfer in the coevolution of the biosphere and
582 geosphere. *Annu. Rev. Microbiol.* **70**.
- 583 79. Jansson JK, Hofmockel KS. 2019 Soil microbiomes and climate change. *Nat. Rev. Microbiol.* , 1–12.
- 584 80. McGenity TJ. 2018 2038 – When microbes rule the Earth. *Environ. Microbiol.* **20**, 4213–4220. (doi:10.1111/1462-2920.14449)
- 585 81. Tornos F *et al.* 2018 Do microbes control the formation of giant copper deposits? *Geology* **47**, 143–146.
- 586 82. Falkowski PG, Fenchel T, Delong EF. 2008 The microbial engines that drive Earth’s biogeochemical cycles. *Science* (80-
587). **320**, 1034–1039.
- 588 83. Schwartzman DW, Volk T. 1989 Biotic enhancement of weathering and the habitability of Earth. *Nature* **340**, 457.
- 589 84. Schwartzman DW. 2017 Life’s critical role in the long-term Carbon cycle: The biotic enhancement of weathering. *AIMS Geosci.*
590 **3**, 216–238.
- 591 85. Bardgett RD, Freeman C, Ostle NJ. 2008 Microbial contributions to climate change through carbon cycle feedbacks. *ISME J.* **2**,
592 805.
- 593 86. Conrad R. 2009 The global methane cycle: Recent advances in understanding the microbial processes involved. *Environ.*
594 *Microbiol. Rep.* **1**, 285–292. (doi:10.1111/j.1758-2229.2009.00038.x)
- 595 87. McCoy DT, Burrows SM, Wood R, Grosvenor DP, Elliott SM, Ma P-L, Rasch PJ, Hartmann DL. 2015 Natural aerosols explain
596 seasonal and spatial patterns of Southern Ocean cloud albedo. *Sci. Adv.* **1**, e1500157.
- 597 88. Sanchez KJ *et al.* 2018 Substantial seasonal contribution of observed biogenic sulfate particles to cloud condensation nuclei. *Sci.*
598 *Rep.* **8**, 1–14.
- 599 89. Carrión O, Curson ARJ, Kumaresan D, Fu Y, Lang AS, Mercadé E, Todd JD. 2015 A novel pathway producing dimethylsulphide
600 in bacteria is widespread in soil environments. *Nat. Commun.* **6**, 1–8.
- 601 90. Jardine K *et al.* 2015 Dimethyl sulfide in the Amazon rain forest. *Global Biogeochem. Cycles* **29**, 19–32.
- 602 91. Bryan NC, Christner BC, Guzik TG, Granger DJ, Stewart MF. 2019 Abundance and survival of microbial aerosols in the
603 troposphere and stratosphere. *ISME J.* **13**, 2789–2799.
- 604 92. Hamilton WD, Lenton TM. 1998 Spora and Gaia: how microbes fly with their clouds. *Ethol. Ecol. Evol.* **10**, 1–16.
- 605 93. Charlson RJ, Lovelock JE, Andreae MO, Warren SG. 1987 Oceanic phytoplankton, atmospheric sulphur, cloud albedo and
606 climate. *Nature* **326**, 655.
- 607 94. Gunson JR, Spall SA, Anderson TR, Jones A, Totterdell IJ, Woodage MJ. 2006 Climate sensitivity to ocean dimethylsulphide
608 emissions. *Geophys. Res. Lett.* **33**.
- 609 95. Gabric AJ, Qu B, Rotstayn L, Shephard JM. 2013 Global simulations of the impact on contemporary climate of a perturbation to
610 the sea-to-air flux of dimethylsulfide. *Aust. Meteorol. Oceanogr. J.* **63**, 365–376.
- 611 96. Wang S, Maltrud ME, Burrows SM, Elliott SM, Cameron-Smith P. 2018 Impacts of shifts in phytoplankton community on clouds
612 and climate via the sulfur cycle. *Global Biogeochem. Cycles* **32**, 1005–1026.
- 613 97. Bak P. 1993 Self-organized criticality and Gaia. In *Thinking about Biology: An Invitation to Current Theoretical Biology* (eds W
614 Stein, F Varela), pp. 255–268. Miami: Santa Fe Institute.
- 615 98. Levin SA. 2005 Self-organization and the emergence of complexity in ecological systems. *AIBS Bull.* **55**, 1075–1079.
- 616 99. Lenton TM, van Oijen M. 2002 Gaia as a complex adaptive system. *Philos. Trans. R. Soc. London B Biol. Sci.* **357**, 683–695.
- 617 100. Schwartzman D. 2002 *Life, temperature, and the Earth: the self-organizing biosphere*. Columbia University Press.
- 618 101. Levin SA. 1998 Ecosystems and the biosphere as complex adaptive systems. *Ecosystems* **1**, 431–436.
- 619 102. Raoult D. 2010 The post-Darwinist rhizome of life. *Lancet* **375**, 104–105.
- 620 103. Morowitz HJ. 1993 *Beginnings of Cellular Life: Metabolism Recapitulates Biogenesis*. New York: Yale University Press.

- 621 104. Morowitz HJ, Smith E, Srinivasan V. 2008 Selfish metabolism. *Complexity* **14**, 7–9. (doi:10.1002/cplx.20258)
- 622 105. Goldford JE, Segrè D. 2018 Modern views of ancient metabolic networks. *Curr. Opin. Syst. Biol.* **8**, 117–124.
623 (doi:10.1016/j.coisb.2018.01.004)
- 624 106. Kim H, Smith HB, Mathis C, Raymond J, Walker SI. 2019 Universal scaling across biochemical networks on Earth. *Sci. Adv.* **5**,
625 eaau0149.
- 626 107. Williams GR. 1996 *The Molecular Biology of Gaia*. New York: Columbia University Press.
- 627 108. Medini D, Donati C, Tettelin H, Massignani V, Rappuoli R. 2005 The microbial pan-genome. *Curr. Opin. Genet. Dev.* **15**, 589–
628 594.
- 629 109. Stolz JF. 2016 Gaia and her microbiome. *FEMS Microbiol. Ecol.* **93**, fiw247.
- 630 110. Magnabosco C *et al.* 2018 The biomass and biodiversity of the continental subsurface. *Nat. Geosci.* **11**, 707–717.
631 (doi:10.1038/s41561-018-0221-6)
- 632 111. Goody RM, Yung YL. 1995 *Atmospheric radiation: theoretical basis*. London: Oxford university press.
- 633 112. Feulner G. 2012 The faint young Sun problem. *Rev. Geophys.* **50**.
- 634 113. Waltham D. 2019 Intrinsic Climate Cooling. *Astrobiology* **19**, 1388–1397.
- 635 114. Owen T, Cess RD, Ramanathan V. 1979 Enhanced CO₂ greenhouse to compensate for reduced solar luminosity on early Earth.
636 *Nature* **277**, 640–642.
- 637 115. Haq-Misra JD, Domagal-Goldman SD, Kasting PJ, Kasting JF. 2008 A revised, hazy methane greenhouse for the Archean Earth.
638 *Astrobiology* **8**, 1127–1137.
- 639 116. Goldblatt C, Matthews AJ, Claire M, Lenton TM, Watson AJ, Zahnle KJ. 2009 There was probably more nitrogen in the Archean
640 atmosphere or This would have helped resolve the Faint Young Sun paradox. *Geochim. Cosmochim. Acta Suppl.* **73**, A446.
- 641 117. Mitchell JFB, Johns TC, Gregory JM, Tett SFB. 1995 Climate response to increasing levels of greenhouse gases and sulphate
642 aerosols. *Nature* **376**, 501–504.
- 643 118. Feichter J, Roeckner E, Lohmann U, Liepert B. 2004 Nonlinear aspects of the climate response to greenhouse gas and aerosol
644 forcing. *J. Clim.* **17**, 2384–2398.
- 645 119. Zaharescu DG, Burghelua CI, Dontsova K, Reinhard CT, Chorover J, Lybrand R. 2020 Biological Weathering in the Terrestrial
646 System: An Evolutionary Perspective. *Biogeochem. Cycles Ecol. Drivers Environ. Impact* , 1–32.
- 647 120. Walker JCG, Hays PB, Kasting JF. 1981 A negative feedback mechanism for the long-term stabilization of Earth's surface
648 temperature. *J. Geophys. Res. Ocean.* **86**, 9776–9782.
- 649 121. Falkowski PG, Isozaki Y. 2008 The story of O₂. *Science (80-.)*. **322**, 540–542.
- 650 122. Olson SL, Reinhard CT, Lyons TW. 2016 Limited role for methane in the mid-Proterozoic greenhouse. *Proc. Natl. Acad. Sci.* **113**,
651 11447–11452.
- 652 123. Boyle RA, Löscher CR. 2019 A biologically driven directional change in susceptibility to global-scale glaciation during the
653 Precambrian-Cambrian transition. *Biorxiv* , 359422.
- 654 124. Garuba OA, Lu J, Liu F, Singh HA. 2018 The active role of the ocean in the temporal evolution of climate sensitivity. *Geophys.*
655 *Res. Lett.* **45**, 306–315.
- 656 125. Zanna L, Khatiwala S, Gregory JM, Ison J, Heimbach P. 2019 Global reconstruction of historical ocean heat storage and transport.
657 *Proc. Natl. Acad. Sci.* **116**, 1126–1131.
- 658 126. Li C, von Storch J-S, Marotzke J. 2013 Deep-ocean heat uptake and equilibrium climate response. *Clim. Dyn.* **40**, 1071–1086.
- 659 127. Rose BEJ, Rayborn L. 2016 The effects of ocean heat uptake on transient climate sensitivity. *Curr. Clim. Chang. Reports* **2**, 190–
660 201.
- 661 128. Resplandy L *et al.* 2018 Quantification of ocean heat uptake from changes in atmospheric O₂ and CO₂ composition. *Nature* **563**,
662 105–108.
- 663 129. England MH *et al.* 2014 Recent intensification of wind-driven circulation in the Pacific and the ongoing warming hiatus. *Nat.*
664 *Clim. Chang.* **4**, 222–227.
- 665 130. Cheng L, Abraham J, Hausfather Z, Trenberth KE. 2019 How fast are the oceans warming? *Science (80-.)*. **363**, 128–129.
- 666 131. Lambeck K, Esat TM, Potter E-K. 2002 Links between climate and sea levels for the past three million years. *Nature* **419**, 199–
667 206.
- 668 132. Brown MS, Munro DR, Feehan CJ, Sweeney C, Ducklow HW, Schofield OM. 2019 Enhanced oceanic CO₂ uptake along the
669 rapidly changing West Antarctic Peninsula. *Nat. Clim. Chang.* **9**, 678–683.
- 670 133. Hinkle GJ. 1996 Marine salinity: Gaian phenomenon? In *Gaia in Action: Science of the Living Earth* (ed P Bunyard), pp. 99–104.

671 Edinburgh: Floris Book.

672 134. Moore JK *et al.* 2018 Sustained climate warming drives declining marine biological productivity. *Science* (80-.). **359**, 1139–1143.

673 135. Schwartzman D, McMenamin M, Volk T. 1993 Did surface temperatures constrain microbial evolution? *Bioscience* **43**, 390–393.

674 136. Wieczynski DJ *et al.* 2019 Climate shapes and shifts functional biodiversity in forests worldwide. *Proc. Natl. Acad. Sci.* **116**, 587–

675 592.

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