

ESD Ideas: Planetary Antifragility: A new dimension in the definition of the Safe Operating Space for Humanity

Oliver López-Corona^{1,2}, Melanie Kolb³, Elvia Ramírez-Carrillo⁴, and Jon Lovett⁵

¹Cátedras CONACyT, CONABIO, CDMX, Mexico

²Centro de Ciencias de la Complejidad (C3), Universidad Nacional Autónoma de México (UNAM), CDMX, Mexico

³Institute of Geography, Universidad Nacional Autónoma de México (UNAM), CDMX, Mexico

⁴Centro de Investigación en Biodiversidad y Conservación, Universidad Autónoma del Estado de Morelos, CDMX, Mexico

⁵School of Geography, University of Leeds, Leeds, LS2 9JT, United Kingdom, Royal Botanic Gardens, Kew, Richmond, Surrey TW9 3AB, United Kingdom

Correspondence: OLC (lopezolverx@ciencias.unam.mx) and ERC (elviarc@otrasenda.org)

Abstract. Combining well-established non-equilibrium thermodynamic principles and a system dynamics approach, we define for the first time the concept of Planetary Antifragility as changes of Fisher information of Earth's entropy production. As a first approximation for entropy production, we propose to use shortwave global albedo anomalies and provide a first quantitative example with data for the July months in the Northern hemisphere from 1982 to 2010. These preliminary results show a net reduction of 47.63% on albedo's Fisher information. This loss of Antifragility implies a compounding problem because human perturbations such as climate or land-use changes are increasing but at the same time, the planet is losing its capacity to respond to them. We discuss the concept of Antifragility in the context of Safe Operating Space for Humanity and Planetary Boundaries. We conclude that not only does interaction among state variables of the boundaries matter, but more importantly the perturbation response capacity dimension is missing. In conclusion we speculate on improving the operational definition for ecosystems.

Copyright statement. Creative Commons Attribution 4.0 License

1 Introduction

The planet Earth hosts a complex self-organising system, famously described in the Gaia Hypothesis of the physicist James Lovelock and biologist Lynn Margulis in a series of papers in the 1970s (Lovelock, 1972; Margulis & Lovelock, 1974; Lovelock & Margulis, 1973). The system consists of dynamic interactions between both inorganic and organic components that the Gaia Hypothesis considered to be self-regulating. Importantly, the organic living component adapts to changing biogeochemical conditions, under thermodynamic constraints, through the process of evolution by natural selection and also changes those conditions as a result of the adaptations (Prigogine et.al. 1972; Lovelock, 2016).

Although the Gaia Hypothesis has been criticised for being teleological, in other words the system 'self-directs' towards conditions best suited to itself, the evolutionary nature of the complex self-organising interactions means that a system without

adaptative benefits is not likely to continue (Kirchner, 1989; Baerlocher, 1990). Examples of beneficial interactions are the widespread symbioses of photosynthetic plants with fungi in mycorrhizae and bacteria in nitrogen fixing nodules for mutual nutrient exchange (Bonfante & Anca, 2009; Bonfante & Genre, 2010;). To take this one step further, cell organelles, such as mitochondria and chloroplasts, are considered to be facultative prokaryotic endosymbionts in eukaryotic cells (Martin et.al. 25 2015).

This cellular development and subsequent evolution of the planet's biodiversity has had far reaching biogeochemical effects. For example, development of vegetation cover changed the composition of the Earth's atmosphere, albedo of the Earth's surface, hydrological cycle and nutrient availability, thereby providing the conditions for further expansion and diversification of life. This complex, dynamic and interacting system continues even if there are substantial external changes, such as the 30 sequence of Croll-Milankovitch orbital cycles considered to be responsible for repeated ice-ages and thought to be important drivers of species diversification and range changes, including expansion of modern humans (Bennett, 1990; Lovelock, 1989).

In this sense the Earth's self-organising system, as described by the Gaia Hypothesis, can be considered to be 'antifragile' in that the response to a potentially catastrophic stressor, even one of the magnitude of shifts in a planet's orbit, leads to continual reorganisation and adaptation that benefits the system as a whole.

35 Set against this, and related to orbital-driven ice ages, is the rise of the modern human teleological culture that takes actions towards a defined end, and that is consequentialist in that the objective if the action is to be achieved by whatever means makes it possible. For example, replacing the Amazon rainforest with grazing land for cattle destined for meat export, or increased agricultural area for commercial livestock feed, is a conscious decision often justified by a need to feed a growing population of humans and bring financial returns where there would otherwise be only forest (Watson, 2004; López-Corona & 40 Magallanes-Guijón, 2020) .

The substantial changes made to the Earth in the past few hundred years have led to concern being expressed about a state shifts in the ability of the planet's complex system to respond. In this article we explore the concept of Planetary Antifragility, following ideas by Michaelian (2015) and Kleidon (2005), as changes of Fisher information of Earth's entropy production using shortwave global albedo anomalies.

45 Ecosystem antifragility differs from concepts such as resilience and robustness because it contains self-organising and adaptive properties in response to external change. In contrast, a resilient ecosystem is one that can recover from deleterious impacts; and a robust ecosystem is one that is able to resist change. Both resilience and robustness can be incorporated in the concept of antifragility as special cases (Equihua et.al. 2020).

In the next section we briefly review the concept of planetary boundaries from an antifragility perspective and present a 50 mathematical definition. We then use satellite measurements of the difference between the incident and outgoing light spectrum of the Earth's albedo as a proxy of planetary scale entropy production.

1.1 Planetary Boundaries

The seminal work "Planetary Boundaries: Exploring the Safe Operating Space for Humanity" (Rockström et al. 2009) identified nine core biogeochemical processes with threshold values for each variable called the Planetary Boundaries (PBs), within

55 which the authors expect that humanity could operate safely. The central idea of their work is that transgressing one or more
 PBs may lead to a catastrophic planetary tipping point most likely incompatible with modern human organization survival.
 Despite the concept of PBs being widely accepted, there are some issues that remain open: (a) As Rockström and coworkers
 (2009) recognize, PBs are not interdependent, as transgressing one may both shift the position of other boundaries and cause
 them to be transgressed and the true threshold configuration of the individual threshold values of the PBs cannot be established
 60 without a metric of the interaction among the whole network of PBs; (b) Rockström and coworkers (2009) say that the impacts
 of transgressing PBs will be a function of the social-ecological resilience, but as has been raised recently (Equihua et al.
 2020), the concept of resilience is a special and limited case of Antifragility; (c) in their work, Rockström et al. (2009) do
 recognize that although not all processes or subsystems on Earth have well-defined thresholds, human actions that undermine
 the resilience of such processes or subsystems can increase the risk that thresholds will also be crossed in other processes, such
 65 as the climate system. In that sense the capacity of the Earth system to respond to perturbation under the resilience concept
 is a key feature in PBs framework. But resilience is a limited special case of the wider concept of Antifragility consisting of
 an intermediate type of response to perturbations near to robustness where the systems tolerate stress and remains the same
 (Equihua et al. 2020).

As a way of emphasizing the importance and rationale of switching from resilience to Antifragility, consider that all living
 70 systems are undergoing evolutionary processes that require them to do far more than simply endure perturbations. Evolutionary
 adaptation requires that they must have some features that allow them to not only cope but gain (up to a point) from pertur-
 bations, stressors, variability, and uncertainty. This is simple evolutionary and natural selection logic: Any system that does
 gain from variability eventually will outperform its competitors over time. As time is the ultimate source of aleatory, many
 different types of phenomena can be expressed through time and an antifragile system will accumulate any gains obtained
 75 through aleatory over time. The consequence of this process is that in contemporary populations and ecosystems we see the
 expression of the accumulation of these advantages over time.

So, through evolutionary processes, what we observe in the present is the predominance of those systems that have previously
 gained from environmental variability and perturbations. This feature is what Taleb calls Antifragility (Danchin et al., 2011;
 Taleb, 2012) and has a formal definition as a non-linear response in the payoffs space and can be summarized as follows (Taleb
 80 and Douady, 2013; Taleb, 2018):

Let $f(x)$ be a two times continuously differentiable payoff function $f(x)$ with a convexity defined as usual by its second
 derivative $\frac{\partial^2 f}{\partial x^2} \geq 0$ which can be simplified without loss of generality as $\frac{1}{2} [f(x + \Delta x) + f(x - \Delta x)] \geq f(x)$.

From this, we can see that as the dose increases there will be a much higher impact (non-linear) response in $f(x)$ which
 generalizes to a linear combination as $\sum \alpha_i = 1, \quad 0 \leq \alpha_i \leq 1$ in such a way that $\sum [\alpha_i f(x_i)] \geq f[\sum (\alpha_i x_i)]$. Then under the
 85 correct conditions we may simplify the argument to $f(nx) \geq nf(x)$, which implies that the payoff function $f(x)$ of the random
 variable X with support in $[a, b]$ we will satisfy the Jensen's Inequality and then

$$\mathbb{E}(f(x)) \geq f(\mathbb{E}(x)). \tag{1}$$

Or as shown in previous work (Taleb and Douady, 2013), the expectation of f under a probability density distribution $\varphi(x)$ with support in $[a, b]$ indexed by the scale σ is

$$90 \quad \forall \sigma_2 > \sigma_1, \mathbb{E}_{\sigma_2}[f(x)] \geq \mathbb{E}_{\sigma_1}[f(x)]; \quad (2)$$

which means that we have either a convex dose-response behavior over $[a, b]$ or the expectation increases with the scale of the distribution.

Given this precise mathematical definition of antifragility, the problem at hand is to identify a suitable systemic payoff
95 function that adequately captures the idea of Planetary Antifragility in the context of an enhanced "Safe Operating Space for Humanity" that incorporates not only PBs' state but also the capacity of the Earth to respond in a convex way to the anthropogenic pressures causing the PBs state to decline.

From previous work on Ecosystem Antifragility (Equihua et.al. 2020) we know that it can be measured using changes in ecosystem complexity and so the most healthy state should be the one that is at level of complexity when the ecosystem can
100 self-organise, a state that is reached when ecosystems are in criticality, a dynamic regime characterized by scale invariance in the Fourier space and also in balance between informational emergence and self-organization (Ramírez-Carrillo et.al. 2018). Finally, we also know that in criticality, systems reach their maximum of Fisher information (López-Corona and Padilla, 2019). In this way, Antifragility may be approximated using the Fisher information as a payoff time series.

The main idea in this context is that the definition of a safe human operating space should consider not only a "safe" range
105 of important state variables in terms of tipping points, but also the dynamics of the system (see for example unpublished work by Toledo-Roy and coworkers <https://www.youtube.com/watch?v=WzfdnoC3Kik>), especially its capacity to respond to perturbations. The dynamic interpretation of Fisher information could be understood as a measure of the system stability or as we have proposed elsewhere (López-Corona and Padilla, 2020) as a universal payoff function for antifragility.

Also there is cumulative evidence presented in our previous work on Ecosystem Antifragility (Equihua et.al. 2020), that
110 points to the conjecture that the Earth System should not only be under a limited range for key biogeochemical variables but also in a criticality regime (Hidalgo et.al. 2014) in which the system is at its maximum complexity, maximum Fisher Information, and balance between emergence (flexibility/randomness) and self-organization. Under these conditions, ecosystems (including Earth systems) exhibit the greatest computational and inferential capacities related to the system capacity to respond and adapt to perturbations.

115 2 Methods

2.1 Entropy production as Payoff function

Albedo anomalies as antifragility estimates

As noticed in previous work by Michaelian (2005, 2015), ecosystems arise and evolve, as any other physical system, under the laws of thermodynamics. In particular, the relation between entropy production and ecosystem functioning up to the Earth system is well established and has been studied since 1972 in the pioneering work of Prigogine and coworkers (1972); then by Ulanowicz and Hannon (1987); Aoki (1989); Schneider and Kay (1994); Schymanski et.al. (2010); Michaelian (2005, 2012, 2015); Kleidon (2005, 2009, 2010a, 2010b, 2010c, 2021); Panwar et.al. (2021).

In the current paper, we build our new Planetary Antifragility concept mainly on the work of Michaelian (2015) who proposes that healthy ecosystems have a bigger entropy production than unhealthy or stressed ones and entropy production should then be a reliable indicator of their health, given by the following expression of total entropy production (more precisely photon dissipation) per unit area of the ecosystem (J) (Michaelian, 2015):

$$Health = J = \int_0^{\infty} 2\pi L_{rad}(\lambda) - 0.04 L_{in}(\lambda) d\lambda, \quad (3)$$

in which the 2π comes from solid angle calculation assuming isotropic emission and predominantly Lambertian reflection as a consequence of leaves diffuse scattering and collective effect of multiple leaves in the ecosystem (Gates, 2012). The 0.04 factor comes from the cosine of the angle of the incident radiation, and both entropy fluxes [$Jm^{-2}K^{-1}$], $L_{rad}(\lambda)$ and $L_{in}(\lambda)$ comes from the following expression as a function of energy flux $I(\lambda)$ [$Jm^{-3}s^{-1}$]. In the case of $L_{rad}(\lambda)$, we have $I = I_{rad}$ measured by a detecting spectrometer; and for $L_{in}(\lambda)$ the $I = I_{in}$ measured by solar spectrum at surface.

$$L(\lambda) = \frac{n_0 k c}{\lambda^4} \left[\left(1 + \frac{\lambda^5 I(\lambda)}{n_0 h c^2} \right) \ln \left(1 + \frac{\lambda^5 I(\lambda)}{n_0 h c^2} \right) - \left(\frac{\lambda^5 I(\lambda)}{n_0 h c^2} \right) \ln \left(\frac{\lambda^5 I(\lambda)}{n_0 h c^2} \right) \right], \quad (4)$$

where λ is the wavelength of energy flux $I(\lambda)$ [$Jm^{-2}s^{-1}$]; c is the velocity of light; n_0 is the photon's polarization state ($n_0 = 1$ for polarized and $n_0 = 2$ for unpolarized); h and k are as usual the Planck and Boltzmann constants.

The key point here is that we don't need to account for all individual living systems that have several non-linear couplings among both biotic and abiotic (i.e. water cycle) components of the Earth system, but can use a global planetary 'good enough' approximation of the entropy production (more precisely photon dissipation) contribution by the biosphere. Nevertheless, it is necessary to find a specific proxy that contains the most relevant information on the entropy production of ecosystems that is measured at a planetary scale and available from existing databases. So, following original ideas discussed by Ulanowicz (1987) and Michaelian (2015), we propose that this proxy can be represented by the difference between the incident and outgoing light spectrum based on satellite measurements of Albedo as in Eq(3).

In this work, we used published data for Surface Albedo anomalies of the Northern hemisphere during the months of July (GLASS albedo product) for 1981–2010 (He et.al., 2014) and calculated its Fisher information in the same way Ahmad and coworkers (2016) have done for global mean temperature (1880-2015). In their work, the authors organized the time series data such that each month represents one system variable so that 12 variables describe global temperature anomalies from January to December for each time step (year).

2.2 Data inputs

Data used here (Fig.1), comes from The Global Land Surface Satellites (GLASS), which uses very high resolution radiometers (AVHRR) and MODIS (Moderate Resolution Imaging Spectroradiometer) data (Liang et.al. 2013). Two direct albedo calculations are incorporated for the MODIS component; one for surface reflectance and one for Top of Atmosphere (TOA) Reflectance (Qu et.al. 2014). The AVHRR observation GLASS albedo component is based on a direct measurement algorithm using radiometric calibration and atmospheric correction surface reflectance (Pedelty et.al. 2007) comparable to that used on MODIS data (Liu et.al. 2013).

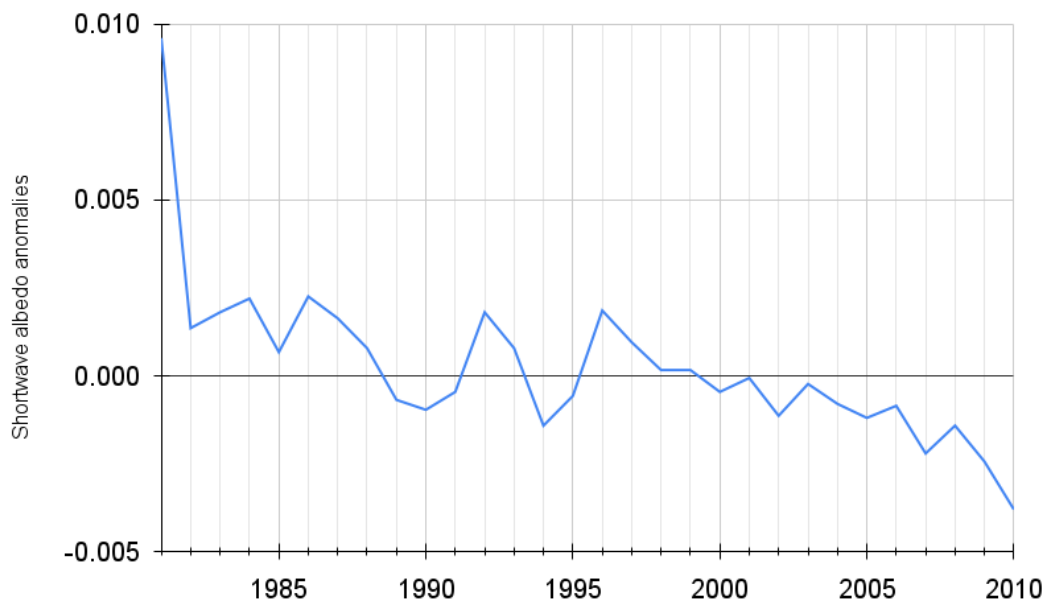


Figure 1. July shortwave terrestrial albedo time series from GLASS data for Northern Hemisphere

We used the 1982 to 2010 time series of July months for the Northern Hemisphere (more land area implies more albedo changes and higher rates of climate change are expected), which show a decrease rate of 0.0013 per decade ($p < 0.01$) (Figure 1)(He et.al. 2014). We took this data set because it is the season when less snow or ice may be found in the Northern Hemisphere and so, a clearer response by the vegetation to Climate Change could be found. We then used a Python implementation of the algorithm created by Ahmad et al. (2016) to assess Fisher information, available at <https://github.com/csunlab/fisher-information>.

In this work, we are only using land albedo, as the original input does not include ocean surface albedo. The different albedo products derived from satellite imagery, in general, are processed only for terrestrial surfaces because ocean albedo is rather stable and low, which may cause higher fluctuations to get masked/averaged by ocean albedo. So, the influence of periodic phenomena like El Niño Southern Oscillation (ENSO) could not affect the results because of the change in ocean surface

albedo. But it must be considered that land cover, which influences terrestrial albedo to a large extent, is heavily influenced by the teleconnections caused by “spatially and temporally large-scale anomalies that influence the variability of the atmospheric circulation” (ENSO, Arctic Oscillation, North Atlantic Oscillation, Pacific Decadal Oscillation, Pacific-North America Index (<https://www.ncdc.noaa.gov/teleconnections/>)). This leads to climate anomalies linked across geographically separated regions, and also to bigger or smaller changes in land cover type (e.g. arid environments with abrupt growth of annual plants after unusual precipitation or drought-related phenotypical changes in normally humid areas). For further work after this proof of concept, it will be interesting to compare the dynamics of both hemispheres.

2.3 Stability analysis using Fisher Information

Consider a dynamical system characterized by a phase space $s = (x_1, \dots, x_i, \dots, x_m)$ built up n state variables x_i under which a measurement y is made, then we can define the quality of the measurement by its Fisher information

$$I(s) = \frac{1}{T} \int_0^T \frac{s''^2}{s'^4} dt, \quad (5)$$

where τ is the time period required for the system to complete a cycle; $s'(t)$ and $s''(t)$ are the tangential velocity and acceleration of the system in the phase space, calculated as a function of the state variables x_i given by

$$s'(t) = \sqrt{\sum_i^m \left(\frac{dx_i}{dt}\right)^2}, \quad (6)$$

$$s''(t) = \frac{1}{s'(t)} \sum_i^m \left(\frac{dx_i}{dt} \frac{d^2x_i}{dt^2}\right). \quad (7)$$

This dynamic interpretation of Fisher information (Frieden et.al. 2007; and Fath, 2002) implies that if the system is a constraint to small tangential velocities and acceleration, then in a specific measurement time range the system will occupy a small hyper-volume, which is interpreted as the system is stable.

In this way, dynamic stability corresponds with higher levels of constant Fisher information. It then follows that self-organized systems would tend to reduce their variability by gaining Fisher information. On the other hand, a loss of stability would be indicated by a reduction in the system’s Fisher information and sudden sharp changes could be used as an early warning of tipping points.

3 Results

In Fig. 2 we show the Fisher Information for 1988–2010 (the algorithm requires an initial calculation using the first 7 data points) of Northern Hemisphere albedo, which exhibits an oscillation with a mean value of 3.59, a maximum value of 5.35 in 1988, minimum of 2.55 in 1998 and a net reduction of 47.63% between 1988 and 2010. The reduction does not occur linearly,

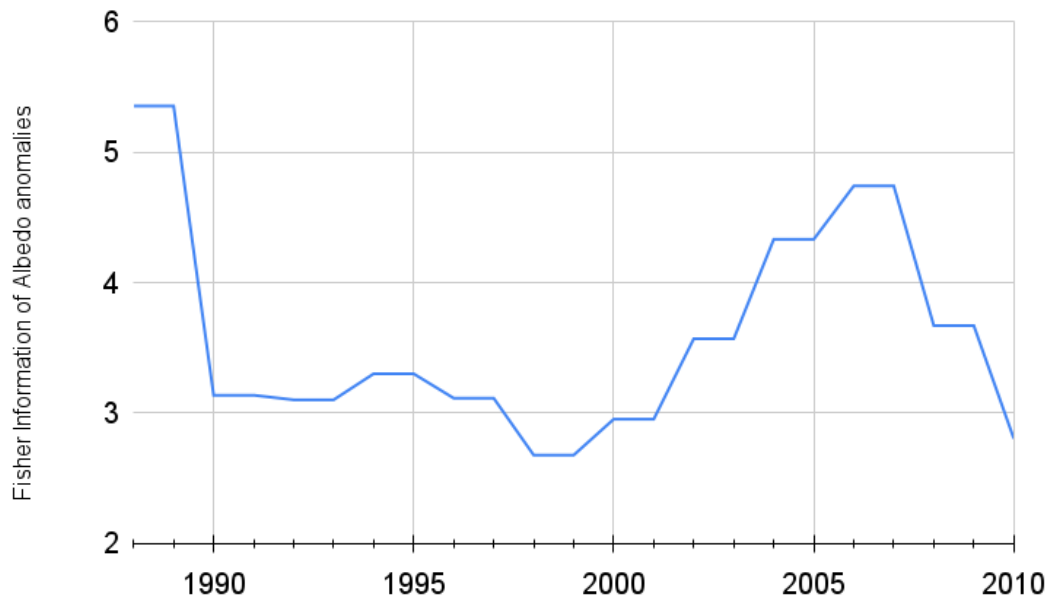


Figure 2. Fisher information for July shortwave terrestrial albedo time series from GLASS data for Northern Hemisphere

190 but shows an oscillating behavior: A first decrease happened between 1988 and 1998, followed by a gradual increase until 2007 to again decrease until 2010.

These results are not meant to provide a complete data analysis on Planetary Antifragility using Fisher information of land albedo (for example, consider the oceanic and hemispheric contributions already discussed). In this paper we are presenting a preliminary analysis to demonstrate that it is possible to evaluate planetary antifragility in a scientifically sound way (Fig. 3) and more importantly, to point the way to further future development of this original idea.

It is important to note that an ESD Ideas article type such as the present one, "presents innovative and well-founded scientific ideas in a concise way that have not been comprehensively explored"; and the main result of the article is the *Planetary Antifragility* idea itself (Fig. 3). This idea opens several paths: (1) a way to use a unified co-evolutionary representation for coupled human and natural systems (CHANS); (2) introduce a precise and quantifiable mathematical framework to systematize the interdisciplinary nature of CHANS; (3) promote and enhance the understanding of the dynamics of emerging, transitional and extreme regimes; (4) increase the interest in developing Artificial Intelligence tools (including Machine or Deep Learning) for interdisciplinary analysis beyond a mechanistic paradigm; (4) the potential to incorporate mathematical methods to improve decision making processes, incorporating CHANS risks aspects.

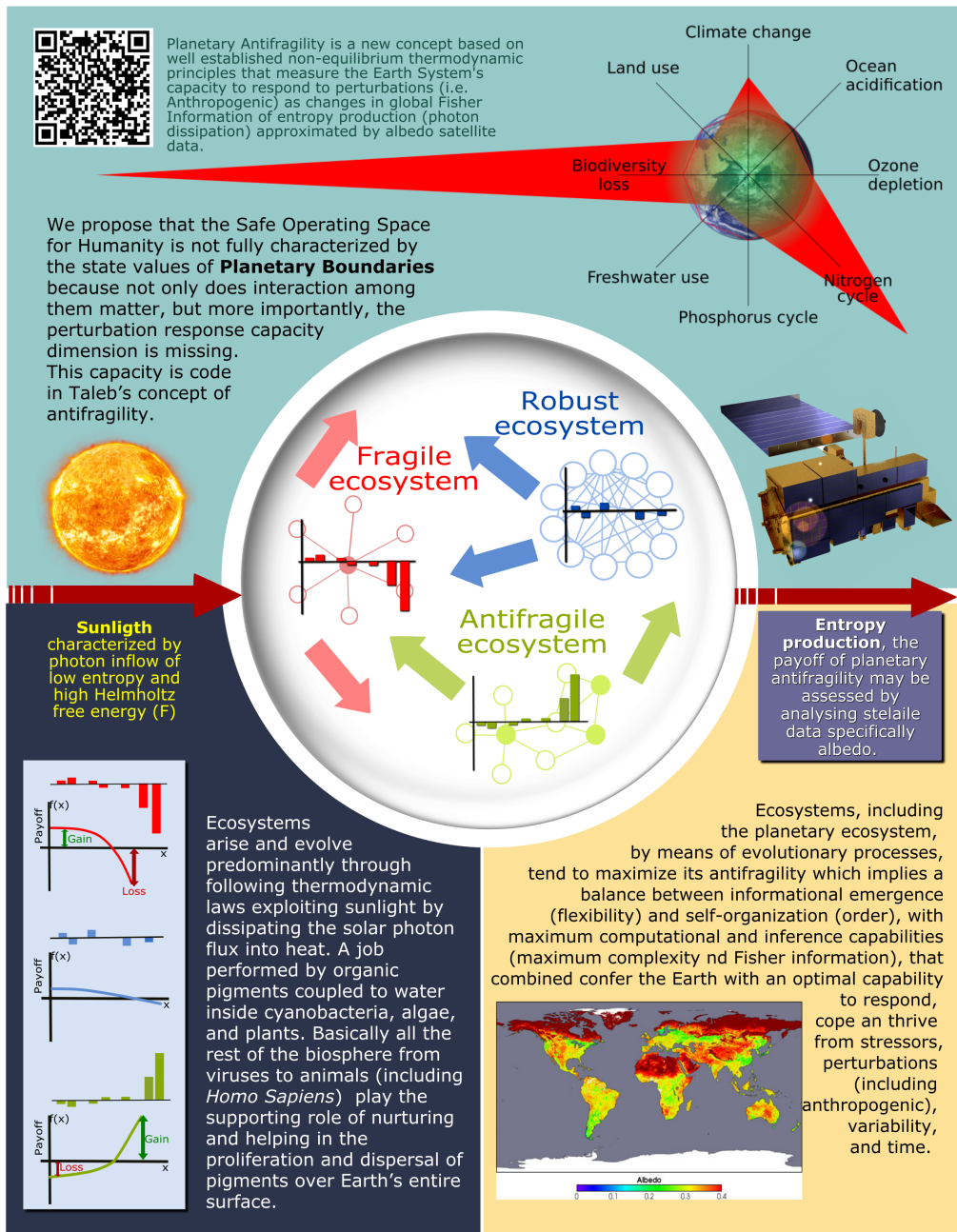


Figure 3. Original figure for the article. Planetary Antifragility is a new concept based on well established non-equilibrium thermodynamic principles that measure the Earth System's capacity to respond to perturbations (i.e. Anthropogenic) as changes in global Fisher Information of entropy production (Photon dissipation) approximated by albedo satellite data.

4 Discussion

205 The "Safe Operating Space for Humanity" is not fully characterized by the state values of Planetary Boundaries because not only does interaction among them matter, but more importantly the perturbation response capacity dimension is missing.

As pointed out in a recent work (Hillebrand et.al. 2020), a key dimension of understanding ecosystem responses to anthropogenic global change is to test if the ecosystem really goes through thresholds or tipping points. In their work, the authors found that threshold transgressions were rarely detectable, either within or across meta-analyses.

210 Has also commented on by Dudley and Suding (2020), and as we have been highlighting in this work, ecosystems seldom respond to environmental drivers in isolation and the inclusion of interacting drivers may indicate more frequent threshold dynamics than expected from meta-analyses (Hillebrand et.al. 2020). In this way, our thermodynamic framework using global albedo as a proxy of planetary entropy production could be interpreted as a systemic response that integrates all drives and responses.

215 The informational approach using Fisher Information as a measure of the entropy production stability leads to the notion of homeostasis, which has been re-framed in terms of time series analysis in a medical application (Fossion et. al. 2018). In their work, they found that when the human body needs to maintain some homeostatic physiological process (keep it within a defined range of values) such as blood pressure. This is achieved by coupling it with another process that absorbs variability from the environment. For homeostasis of blood pressure, the heart rate needs to absorb environmental volatility. They show
220 that healthy people have a blood pressure that is normally distributed, while heart rate is skewed to the right. Whereas when there is a chronic disease such as diabetes, blood pressure is no longer Gaussian and generates a skew to the left, while heart rate becomes normally distributed. When a process is normally distributed, it means that there is a well-defined characteristic scale around which all values are clustered, with very few extreme values. Conversely, having skewed distributions means that there are many extreme events, which in fact dominate the phenomenon to the degree that the characteristic scale can be lost.

225 Homeostatic processes could explain why threshold transgressions were rarely detectable in ecosystems that are self-organising to prevent tipping point events. For example, we could consider ecosystem functions as homeostatic processes maintained by the fluctuations in species composition. In this case, compositional shifts should be much more prone to threshold dynamics than ecosystem functions.

Also interesting is the similarity between the results of Fossion and coworkers (2018) with Taleb's ideas about Antifragility
230 (Taleb, 2012). This made us reconceptualize ecosystem homeostasis or resilience, as it is generally identified in ecology, as a particular case of Taleb's conceptual framework, in which a system can be fragile, robust, or antifragile, depending on how it responds to disturbances in its environment (see Fig. 1). As quantitative Antifragility is measured as the system's response to perturbation, in order to evaluate it one needs to identify the adequate payoff function; in this case, we argue that planetary entropy production can be approximated by Earth's albedo.

235 Using published data (He et.al. 2014) we calculated Fisher Information for 1988–2010 of Northern Hemisphere albedo, which exhibits an oscillation with a mean value of 3.59, a maximum value of 5.35, minimum of 2.55, and a net reduction of 47.63%; with a rate of 0.0013 per decade ($p < 0.01$) that is in agreement with findings by (Marcianesi et. al. 2020) who

calculated, for example, a global Clear-sky albedo (%) decrease of 0.24% per decade, with the confidence of 99% for land; and a decrease of 0.66% per decade, with the confidence of 99% for land. In this context land-use changes might be an important driver for Antifragility loss.

Observed oscillation in Fisher information values could be interpreted as a cyclical decrease in Fisher information, as the increase after the completion of one cycle does not rebound to the original value but stays below. This decrease would be associated with a loss of stability (degradation?) overlapped with oscillations caused by changes in terrestrial albedo as a response to teleconnected climate oscillations.

Another interpretation related to this cyclical pattern could be based on critical slowing down: Strogatz (1994) proposed critical slowing down as representing the major contribution from the authors. Critical slowing down “implies that recovery upon small perturbations becomes slower as a system approaches a tipping point” (Scheffer et al. 2015). This could explain why the recovery after the first cycle of loss of Fisher Information does not reach the original value, as a slowing down means less recovery in the same amount of time. If the time of recovery and loss of Fisher information is determined by oscillating climate phenomena, a slowing down of recovery would mean less recovery between cycles.

But considering the overall Fisher Information lost, it is not only that the planet is decreasing its albedo as a response to human perturbations (mainly climate change and land use change) as the data show: If albedo is a proxy of entropy production, what Michaelian recognized as the thermodynamic function of life (Michaelian, 2012), then loss of albedo’s stability means the planet is losing a key feature of its dynamics: in other words, it’s Antifragility.

This means that we may have a compounding problem because human perturbation such as climate change is increasing, but at the same time the planet is losing its capacity to respond to it. In that sense, we need to not only reduce or capture CO2 emissions, but we should also restore Earth’s Antifragility, which means restoring its ecosystems.

Now let us consider some problems or limitations: (1) In human health assessment the first-order approximation has been identified with the previously known reference range of value of some key physiological variables such as heart rate and systolic blood pressure. So one problem with using visible albedo as a proxy for global entropy production is we do not have the equivalent of those reference values, which in this case should be determined for each ecosystem type. In that sense, visible albedo should be applied in a spatially explicit way, not averaging mean values over large regions, but using local values because the values need to be evaluated against the correct reference values.

(2) Another problem would be considering visible albedo values without their dynamics, as can be illustrated with an analogy to human health: Consider a person with a broken arm (unhealthy state) but a healthy heart (healthy dynamics) versus an Olympic athlete (healthy state) but with a condition prone to sudden cardiac syndrome (unhealthy dynamics). Given these two considerations, we decided not to rely on the direct value of albedo but rather its Fisher information, which encodes the system’s dynamics in terms of its capacity to respond to perturbations.

(3) As suggested by Michaelian (2015), other problems could be that the real extent of the ecosystem considered in the measurement depends on the height of the remote sensor because of the relation with the solid angle of the detector.

(4) Consider that an albedo value is an “instant” measurement and it could be necessary to integrate measurements across a 24 h cycle (if possible) or other longer cycles; but perhaps this is not important for a long term analysis, as presented in this work.

Nevertheless, it does point to the fact that this work does not present a fully developed framework for Planetary Antifragility, we still need to resolve if remote sensing measurements of albedo really can sufficiently capture entropy production, or if
275 other signals would be needed. This would most likely be the case when working at a more detailed scale, for example, on a particular ecosystem in a defined region.

This brief concept paper is a first illustration of the general idea of using antifragility as a new dimension in the definition of the "Safe Operating Space for Humanity". As a follow-on from discussing some aspects of the advantages and problems with using albedo measurements derived from satellite imagery in this manuscript, we would very much like to further explore
280 other variables that could be used to construct indicators for planetary antifragility.

In addition to albedo, we think it would be very interesting to eventually incorporate other measurable proxies for entropy production sources, such as the bioacoustic signals and perhaps also ecosystem respiration. Every sound emitted by a living agent in an ecosystem is coding a part of the ecosystem metabolism into the signal. Important bioacoustic signals are produced by members of the animal kingdom and would prevent the problem posed by defaunated ecosystems, which
285 from a vegetation perspective using measures such as albedo, could give a signal of being healthy (at least in the short term). Ecosystem respiration is a good proxy because it includes soil respiration, and soil is a complex system that incorporates all spheres (biosphere, geosphere, atmosphere, hydrosphere), several biogeochemical processes, many spatial and time scales, so it links many sources of information about the ecosystem in a single measure. Although there are major projects as Ameriflux (<https://ameriflux.lbl.gov/>) that collect data of ecosystem respiration (among others) in many places of America (mostly North
290 America), currently there is not enough data available for considering these variables at a planetary scale. However the concept may downscale to ecosystem-scale using both remote sensing and in situ data, which could work also as a benchmark against other ecosystem health metrics such as integrity.

Finally, as always happens when introducing a new concepts, the revised perspective leads us to revisit others. In this case, the ecosystem definition that traditionally is understood in its basic form as a spatially explicit community of living organisms
295 in conjunction with the nonliving components of their environment, and which interacts as a system. As pointed out by Jax (2007) managing ecosystems or increase their theoretical understanding, requires a clear conceptualization of what ecosystems are (Schaeffer and Cox 1992; Sagoff 2003; Jax 2005); however, there are different and sometimes incompatible definitions of the notion of ecosystem (see e.g. Jax et al. 1998; Jax 2006).

Depending on the specific definition under use, there might be different sets of questions that are difficult to answer. In
300 particular, under the basic definition given above: How many different species and interactions are necessary to constitute an ecosystem? Is a community of trees of different species an ecosystem? What about function, dynamics, stability? Are there any requirements in terms of spatial extension and connectivity? Are there different types of ecosystems for example, what happens with modified, degraded, or perturbed "ecosystems"? To what extent is a "modified" ecosystem of type A, still an A ecosystem? Based on what characteristics a type of ecosystem should be defined? What happens with different ecosystems
305 that share some interactions, species? Does this definition allow us to understand predicted succession, and ecological tipping points?

Also, as commented by Sagoff (2003), to improve our understanding about ecosystems, including the planetary, its definition should: (1) define and classify them, and thus determine the conditions under which it remains the "same" system through time and change; (2) find ways to reject as well as to create mathematical models of the ecosystem functioning and evolution; (3) 310 make possible to identify efficiently the causes of ecosystem organization; (4) show how ecology can potentially help to solve socio-environmental problems.

We think that the Antifragility framework could help to improve ecosystem definition in a manner that many of these questions and goals are easier to tackle. So we are incorporating the ideas of this paper in this new definition: An ecosystem is an open thermodynamic system constituted by a community of living organisms in conjunction with the nonliving components 315 of their environment; that through its interactions and evolutionary processes, constrained by the external conditions; self-organized in a maximum solar photon flux dissipation, in which the system is at criticality, with maximum computational and inferential capabilities that allows it to respond and thrive under uncertainty, stressors, perturbations and ultimately time, in a well defined geographic context.

Code/Data availability

320 Code available at: <https://github.com/csunlab/fisher-information>

Data is published and available in the corresponding reference (He et.al. 2014).

Author contribution

OL-C contributions: Conceptualization, Data curation, Formal analysis, Investigation, Methodology, Supervision, Writing – original draft preparation, Writing – review and editing

325 MK contributions: Conceptualization, Investigation, Methodology, Supervision, Writing – original draft preparation, Writing – review and editing

ERC contributions: Major revision writing, editing.

JL contributions: Major revision writing, editing.

Competing interests

330 The authors declare that they have no conflict of interest.

Acknowledgments

This work was supported by CONACyT fund M0037-2018-07, Number 296842, Cátedras CONACyT fellowship program (Project Number 30), and Sistema Nacional de Investigadores SNI support. The funders had no role in study design, data

collection and analysis, decision to publish, or preparation of the manuscript. We also thank Jonathan Kenny for valuable
335 proofreading.

5 References

- Aoki, I. (1989). Holological study of lakes from an entropy viewpoint-lake Mendota. *Ecological Modelling*, 45(2), 81-93.
- Ahmad, N., Derrible, S., Eason, T., and Cabezas, H.: Using Fisher information to track stability in multivariate systems, 3,
160582, 2016. *Royal Society Open Science*, 3(11)
- 340 Baerlocher, F. (1990). The Gaia hypothesis: a fruitful fallacy?. *Experientia*, 46(3), 232-238.
- Bennett, K. D. (1990). Milankovitch cycles and their effects on species in ecological and evolutionary time. *Paleobiology*,
16(1), 11-21.
- Bonfante, P., & Anca, I. A. (2009). Plants, mycorrhizal fungi, and bacteria: a network of interactions. *Annual review of
microbiology*, 63(1), 363-383.
- 345 Bonfante, P., & Genre, A. (2010). Mechanisms underlying beneficial plant–fungus interactions in mycorrhizal symbiosis.
Nature communications, 1(1), 1-11.
- Danchin, A., Binder, P. M., and Noria, S.: Antifragility and tinkering in biology (and in business) flexibility provides an
efficient epigenetic way to manage risk, 2, 998–1016, 2011. *Genes* 2(4): 998-1016, DOI: 10.3390/genes2040998
- Dudney, J. and Suding, K. N.: The elusive search for tipping points, 4, 1449–1450, 2020. *Nature Ecology & Evolution* 4:
350 1449-1450, DOI: 10.1038/s41559-020-1273-8
- Equihua, M., Aldama, M. E., Gershenson, C., López-Corona, O., Munguía, M., Pérez-Maqueo, O., and Ramírez-Carrillo, E.:
Ecosystem antifragility: beyond integrity and resilience, 8, e8533, 2020. *PeerJ* 8: e8533. doi: 10.7717/peerj.8533
- Fossion, R., Rivera, A. L., and Estañol, B.: A physicist’s view of homeostasis: how time series of continuous monitoring
reflect the function of physiological variables in regulatory mechanisms, 39, 084007, 2018. *Physiological Measurement*.
355 39 084007, DOI:10.1088/1361-6579/aad8db
- Frieden, B. R.: Introduction to Fisher information: its origin, uses, and predictions, in: *Exploratory data analysis using Fisher
information*, Springer, 1–41, 2007.
- Gates, D. M. (2012). *Biophysical ecology*. Courier Corporation.
- He, T., Liang, S., and Song, D.-X.: Analysis of global land surface albedo climatology and spatial-temporal variation during
360 1981–2010 from multiple satellite products, 119, 10–281, 2014. *JGR Atmospheres* 119(17): 10281-10298.
<https://doi.org/10.1002/2014JD021667>

- Hidalgo, J., Grilli, J., Suweis, S., Munoz, M. A., Banavar, J. R., and Maritan, A.: Information-based fitness and the emergence of criticality in living systems, 111, 10095–10100, 2014. PNAS 111(28): 0095-10100.
<https://doi.org/10.1073/pnas.1319166111>
- 365 Hillebrand, H., Donohue, I., Harpole, W. S., Hodapp, D., Kucera, M., Lewandowska, A. M., Merder, J., Montoya, J. M., and Freund, J. A.: Thresholds for ecological responses to global change do not emerge from empirical data, 4, 1502–1509, 2020. *Nature ecology evolution* 4: 1502-1509.
<https://doi.org/10.1038/s41559-020-1256-9>
- Jax, K., Jones, C. G., & Pickett, S. T. (1998). The self-identity of ecological units. *Oikos*, 253-264. *Oikos* 82(2): 253-264.
370 <https://doi.org/10.2307/3546965>
- Jax, K. (2005). Function and “functioning” in ecology: what does it mean?. *Oikos*, 111(3), 641-648.
<https://doi.org/10.1111/j.1600-0706.2005.13851.x>
- Jax, K. (2006). Ecological units: definitions and application. *The quarterly review of biology*, 81(3), 237-258.
<https://doi.org/10.1086/506237>
- 375 Kirchner, J. W. (1989). The Gaia hypothesis: Can it be tested?. *Reviews of Geophysics*, 27(2), 223-235.
- Kleidon A, Lorenz RD (2005) Entropy production by earth system processes. In: Kleidon A, Lorenz RD (eds) *Non-equilibrium thermodynamics and the production of entropy: life, earth, and beyond*. Springer, Heidelberg. ISBN 3-540-22495-5
- Kleidon, A. (2009). Maximum entropy production and general trends in biospheric evolution. *Paleontological Journal*, 43(8), 980-985.
380 <https://doi.org/10.1134/S0031030109080164>
- Kleidon, A., Malhi, Y., & Cox, P. M. (2010a). Maximum entropy production in environmental and ecological systems. *Philosophical Transactions of the Royal Society B* 365(1545).
<https://doi.org/10.1098/rstb.2010.0018>
- Kleidon, A. (2010b). Life, hierarchy, and the thermodynamic machinery of planet Earth. *Physics of life reviews*, 7(4), 424-
385 460.
<https://doi.org/10.1016/j.plrev.2010.10.002>
- Kleidon, A. (2010c). Non-equilibrium thermodynamics, maximum entropy production and Earth-system evolution. *Philosophical Transactions of the Royal Society A: Mathematical, Physical and Engineering Sciences*, 368(1910), 181-196.
<https://doi.org/10.1098/rsta.2009.0188>
- 390 Kleidon, A. (2021). What limits photosynthesis? Identifying the thermodynamic constraints of the terrestrial biosphere within the Earth system. *Biochimica et Biophysica Acta (BBA)-Bioenergetics*, 1862(1), 148303.
<https://doi.org/10.1016/j.bbabi.2020.148303>

- 395 Liang, S., Zhao, X., Liu, S., Yuan, W., Cheng, X., Xiao, Z., Zhang, X., Liu, Q., Cheng, J., Tang, H., and Others: A long-term Global LAnd Surface Satellite (GLASS) data-set for environmental studies, 6, 5–33, 2013. *International Journal of Digital Earth* 6(1).
<https://doi.org/10.1080/17538947.2013.805262>
- Liu, Q., Wang, L., Qu, Y., Liu, N., Liu, S., Tang, H., and Liang, S.: Preliminary evaluation of the long-term GLASS albedo product, 6, 69–95, 2013. *International Journal of Digital Earth* 6(1).
<https://doi.org/10.1080/17538947.2013.804601>
- 400 López-Corona, O., & Magallanes-Guijón, G. (2020). It is not an Anthropocene; it is really the Technocene: names matter in decision making under planetary crisis. *Frontiers in Ecology and Evolution*, 8, 214.
- López-Corona, O. and Padilla, P.: Fisher information as unifying concept for criticality and antifragility, a primer hypothesis, 2019. DOI: 10.13140/RG.2.2.28789.73444
- Lovelock, J. E. (1989). Geophysiology, the science of Gaia. *Reviews of Geophysics*, 27(2), 215-222.
- 405 Lovelock, J. E., Gaia as seen through the atmosphere, *Atmos. Environ.*, 6, 579-580, 1972.
- Lovelock, J. Gaia: A new look at life on earth. Oxford University Press, 2016.
- Lovelock, J. E., & Margulis, L. Atmospheric homeostasis by and for the biosphere: the Gaia hypothesis. *Tellus*, 26(1-2), 2-10, 1974.
- Marcianesi, F., Aulicino, G., and Wadhams, P.: Arctic sea ice and snow cover albedo variability and trends during the last
410 three decades, 100617, 2020. *Polar Science* 28:100617.
<https://doi.org/10.1016/j.polar.2020.100617>
- Margulis, L., and J. E. Lovelock, Biological modulation of the Earth's atmosphere, *Icarus*, 21, 471-489, 1974
- Martin, W. F., Garg, S., & Zimorski, V. (2015). Endosymbiotic theories for eukaryote origin. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 370(1678), 20140330.
- 415 Michaelian, K.: Thermodynamic stability of ecosystems, 237, 323–335, 2005. *J Theor Biol.* 2005 Dec 7;237(3):323-35.
doi: 10.1016/j.jtbi.2005.04.019.
- Michaelian, K.: HESS Opinions“ Biological catalysis of the hydrological cycle: life's thermodynamic function,” 16, 2629–2645, 2012. *Hydrology and Earth System Sciences* 16, 2629–2645.
<https://doi.org/10.5194/hess-16-2629-2012>
- 420 Michaelian, K.: Photon Dissipation Rates as an Indicator of Ecosystem Health, in: *Environmental Indicators*, Springer, 15–36, 2015. In: Armon, R., Hanninen, O. (eds) *Environmental Indicators*. Springer, Dordrecht.
https://doi.org/10.1007/978-94-017-9499-2_2

- Panwar, A., Kleidon, A., & Renner, M. (2020, December). What Cools Forests: Evaporation or Aerodynamic conductance?. In AGU Fall Meeting Abstracts (Vol. 2020, pp. H192-05).
- 425 Pedelty, J., Devadiga, S., Masuoka, E., Brown, M., Pinzon, J., Tucker, C., Vermote, E., Prince, S., Nagol, J., Justice, C., and Others: Generating a long-term land data record from the AVHRR and MODIS instruments, in: 2007 IEEE International Geoscience and Remote Sensing Symposium, 1021–1025, 2007. 2007 IEEE International Geoscience and Remote Sensing Symposium, 2007, pp. 1021-1025, doi: 10.1109/IGARSS.2007.4422974.
- 430 Prigogine I, Nicolis G, Babloyantz A (1972) Thermodynamics of evolution (I). *Phys Today* 25:23–28; Thermodynamics of evolution (II). *Phys Today* 25 (11):38–44 (1972).
<https://doi.org/10.1063/1.3071090>
- Qu, B., Xi, Z., and Chen, C.: A method for TOA location based on scattering models in NLOS environments, 39, 27–30, 2014. in *IEEE Transactions on Vehicular Technology*, vol. 56, no. 2, pp. 583-593, March 2007,
435 doi: 10.1109/TVT.2007.891491.
- Ramírez-Carrillo, E., López-Corona, O., Toledo-Roy, J. C., Lovett, J. C., de León-González, F., Osorio-Olvera, L., Equihua, J., Robredo, E., Frank, A., Dirzo, R., and Others: Assessing sustainability in North America’s ecosystems using criticality and information theory, 13, e0200382, 2018. *PLOS ONE*
<https://doi.org/10.1371/journal.pone.0200382>
- 440 Rockström, J., Steffen, W., Noone, K., Persson, A.A.S., Chapin, F. S., Lambin, E. F., Lenton, T. M., Scheffer, M., Folke, C., Schellnhuber, H. J., and Others: A safe operating space for humanity, 461, 472–475, 2009. *Nature* 461, 472–475 (2009).
<https://doi.org/10.1038/461472a>
- Sagoff, M. (2003). The plaza and the pendulum: Two concepts of ecological science. *Biology and Philosophy*, 18(4), 529-552. *Biology & Philosophy* 18, 529–552 (2003). <https://doi.org/10.1023/A:1025566804906>
- 445 Schaeffer, D. J., & Cox, D. K. (1992). Establishing ecosystem threshold criteria. *Ecosystem Health-New Goals for Environmental Management*. Island, Washington DC.
- Schneider ED, Kay JJ (1994) Life as a manifestation of the second law of thermodynamics. *Math Comput Model* 19(6–8):25–48.
[https://doi.org/10.1016/0895-7177\(94\)90188-0](https://doi.org/10.1016/0895-7177(94)90188-0)
- Schymanski, S. J., Kleidon, A., Stieglitz, M., & Narula, J. (2010). Maximum entropy production allows a simple representation of heterogeneity in semiarid ecosystems. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 365(1545), 1449-1455.
450 <https://doi.org/10.1098/rstb.2009.0309>

- Strogatz, S. H. (1994). Norbert Wiener's brain waves. In *Frontiers in mathematical biology* (pp. 122-138). Springer, Berlin, Heidelberg.
- 455 Taleb, N. N.: *Antifragile: Things that gain from disorder*, Random House Incorporated, 2012.
- Taleb, N. N.: (Anti) Fragility and Convex Responses in Medicine, in: *International Conference on Complex Systems*, 299–325, 2018.
- Taleb, N. N. and Douady, R.: Mathematical definition, mapping, and detection of (anti) fragility, 13, 1677–1689, 2013.
<https://doi.org/10.48550/arXiv.1208.1189>
- 460 Ulanowicz, R. E. and Hannon, B. M.: Life and the production of entropy, 232, 181–192, 1987. *Proceedings of the Royal Society of London. Series B, Biological Sciences*, 232(1267), 181–192. <http://www.jstor.org/stable/36217>
- Watson, A. J. (2004). Gaia and observer self-selection. *Scientists Debate Gaia*, 201-208.