## **Open-Ended Evolution: The Cooperation Barrier**

John E. Stewart

Evolution, Complexity and Cognition Group, Center Leo Apostel, Vrije Universiteit Brussel, Krijgskundestraat 33, B-1160 Brussels, Belgium. future.evolution@gmail.com

### **Extended Abstract**

This abstract: (a) identifies a key barrier to open-ended evolution that is encountered each time new levels of organization emerge in the evolution of life; (b) describes the mechanisms that allow the barrier to be overcome (with particular attention given to how the barrier that separates chemical organization from living organization is overcome); and (c) discusses the implications of the barrier for the design and cultivation of artificial life systems.

### The Cooperation Barrier and the Origin of Life

In order for life to emerge from chemical processes, a major barrier to open-ended evolution has to be overcome. This 'cooperation barrier' arises because self-producing organizations of molecular species are able to explore only a very restricted possibility space (Bagley and Farmer, 1991; Kauffman, 1993; Maynard Smith, 1979; Nighe, et al. 2015; and Vasas, et al. 2012).

The nature of this barrier can be understood by considering an organization of molecular species that is self-producing because it is collectively autocatalytic-i.e. the formation of every species in the organization is catalysed by at least one other species, and the organization has access to appropriate sources of free energy and 'food' molecules. The cooperation barrier arises because molecular species that could contribute to the survivability of the organization may not be produced and sustained at an optimal level within the organization. This can be the case irrespective of the significance of the contribution that these species could make to the success of the organization. How can this occur? First, the formation of a particular molecular species might not happen to be catalysed by any other member of the organization (or it may not be catalysed at a level that is optimal for the organization). This is not likely to be uncommon—there is nothing at all in the nature of autocatalytic organization that guarantees that any particular molecular species that contributes to the organization will be catalysed in return. Second, it might occur where 'parasites' and other 'free-rider' molecular species take resources from the organization but do not contribute anything (or insufficient) in return (e.g. they do not catalyse the formation of other members of the organization). Free-riders can reduce the catalytic support, energy and material resources that might otherwise be available to other members of the organization, undermining their ability to persist and contribute to the organization.

Because free-riders do not use their resources to contribute to the organization, they may also out-compete those that do. The susceptibility of an organization to undermining by free-riders is likely to increase as its complexity increases.

If a molecular species is not produced within an organization, the organization containing that species cannot be sustained or called into existence by selection, no matter how powerful the selection is. This seriously limits the extent of the possibility space that can be explored by collectively autocatalytic organizations of molecular species. It also seriously limits the extent of novelty that can arise amongst these organizations and be the subject of selection. As has been noted, the production of sufficient novelty is a prerequisite for open-ended evolution (Taylor, 2015).

# The Cooperation Barrier and Other Major Evolutionary Transitions

The cooperation barrier does not only restrict the space of possible cooperative organizations that can be explored by molecular organizations. It is also a barrier to the emergence of complex cooperative organizations at each and every level of living organization. It is therefore a barrier to the emergence of new levels of organization (and to the open-ended evolution that is facilitated by the emergence of new levels). For example, the cooperation barrier impeded the emergence of the cooperative organizations of eukaryote cells that became multicellular organisms, the organisations of organisms that became animal societies, the organisations of humans that became tribal societies, the organizations of human groups that became nation states, and is currently impeding the emergence of a complex, cooperative planetary entity (it should be noted that these emergences include many but not all of the major evolutionary transitions identified by Maynard Smith and Szathmary (1995) [e.g. it does not include sexual reproduction], and includes emergences that they do not include [e.g. the emergence of a cooperative global organization (Stewart, 2014)]).

A generalized agent-based approach can be used to understand the causes of the cooperation barrier that impedes the exploration of complex organization at every level of organization (Stewart, 2015). Using this approach, agents represent the entities at each particular level (e.g. prokaryote cells, eukaryotes, multicellular organisms [including humans], tribes, nations etc.). Agents are capable of adaptation. Adaptation tends to maximize a function such as fitness or psychological utility. Entitles may adapt by any process (e.g. including by processes as disparate as gene-based natural selection or psychological mechanisms). Agents are able to interact with each other in ways that impact on the success of their adaptations. Cooperative organizations of agents will emerge where adaptations that constitute cooperative relationships between agents are beneficial to the agents (i.e. where the adaptations provide net fitness or utility benefits to the individual agents that exhibit them). Where this condition is met, the relationships and the organization they constitute will persist and be reproduced through time.

However, in many circumstances this condition will not be met, despite the potential of many forms of cooperation to significantly increase the net benefits available to an organization. It will not be met if co-operators fail to capture enough of the benefits they produce to outweigh the costs of their cooperation. As the huge body of research on cooperation referred to below has demonstrated, this failure can be expected to be commonplace. There is nothing in simple, unstructured forms of organization which guarantees that co-operator agents will always capture sufficient of the benefits they create. To the contrary, agents that support co-operators will tend to be outcompeted by agents that use resources only for their own benefit, without providing sufficient benefits to the organization in return (e.g. free-rider agents, including parasites, cheats and thieves). Free-riders will also tend to outcompete the co-operator agents themselves, and take resources that might otherwise support co-operators. Furthermore, there is nothing that guarantees that free-rider agents will always capture the 'harms' that they visit on the organization. For all these reasons, free-rider agents will tend to undermine complex cooperative organization.

As a consequence, the cooperation barrier will seriously restrict the possibility space of complex cooperative organization that can be explored at any level of organization. All forms of organization that include agents that provide significant net benefits to the organization but fail to capture sufficient of those benefits will not be able to persist. These forms of organization will not be able to be produced as part of the extensive novelty that is essential if evolution is to be openended.

### Mechanisms that can Overcome the Barrier

A huge literature exists that attempts to identify particular mechanisms which enable co-operator agents to capture sufficient of the benefits they create to enable the emergence of some form of cooperative organization (e.g. see Stewart, 2014). These mechanisms generally rely on co-operators capturing a disproportionate share of the benefits of cooperation because of the existence of circumstances which ensure they are disproportionately likely to interact with other co-operators. These biased patterns of interaction are typically produced by constraints that manifest as, for example: particular dispersal patterns; kin selection; group formation; compartmentalization; stochastic correction; other forms of population structure; predispositions to cooperate preferentially with other co-operators; and pre-dispositions to punish and exclude free-riders. However, in general this body of research confirms the reality of the cooperation barrier. It has demonstrated that complex cooperative organization does not evolve readily. It has shown that simple cooperative relationships can emerge, but only in limited circumstances. Most researchers in this field would accept that the research has so far been unable to identify a general mechanism that could operate at all levels of organization and that would enable complex cooperative organization to emerge readily.

But the cooperation barrier has been overcome repeatedly and comprehensively during the evolution of life on this planet, enabling the emergence of complex cooperative organization at various levels. What mechanism(s) have enabled this? It is clear from the agent-based perspective sketched above that agents who provide significant net benefits to an organization would be able to persist if 'consequence-capture' applies—i.e. if agents capture sufficient of the benefits (and harms) they produce to sustain them at an optimal level in the organization. Comprehensive consequence-capture would massively expand the possibility space that can be explored by organizations at any level (Stewart, 2015).

But what can produce consequence-capture? 'Managers' can enable comprehensive consequence-capture within the organizations they manage (Stewart, 1995, 1997, 2000, 2014 and 2015). Managers are powerful, evolvable agents (or coalitions of agents) that can control an organization to support co-operators and to suppress free riders. Managers control an organization by applying constraints (Salthe, 1985). Constraints can influence the dynamical behaviour within the organization without being influenced in return (this is the essence of control). Constraints can operate to direct resources preferentially to co-operator agents, and can punish or suppress free-riders. In order to apply constraints, managers must function independently of the dynamical interactions within the organization proper. They must be able to stand outside and be able to act across the dynamic. Managers do not depend on participation in the dynamical interactions within the organization to survive and persist. They can use their constraining power to appropriate whatever resources they need from the organization. Without the capacity to constrain (to influence without being influenced in return), any attempt by managers to appropriate resources for themselves or to distribute resources to particular agents could be undermined by other agents, and free riders could escape control. Just as normal, powerless members of a human organization are unable to control or manage the organization, normal agents within an organization cannot apply constraints to it or begin to manage it-they cannot influence without being influenced in return. The dynamical separation of managers from the organization often results from the fact that the processes that constitute managers are larger in scale, involve slower rate processes and/or are relatively more stable than the processes that constitute the organization proper (Salthe, 1985).

Management and the constraints it applies can be more or less enabling or more or less prescriptive. Where management itself is comprised of a coalition of agents, it will encounter its own cooperation barrier. This barrier can be overcome by constraints that suppress competition within management. Management can be external to the agents that are being managed, or can be internal to the agents and distributed across them. Examples of 'external management' include: RNA/DNA management of a cell; and management of a human society by its government. Examples of 'distributed internal management' include: a multi-cellular organism in which the behaviour of cells is controlled across the organisation by genetic constraints that are reproduced in each cell; and a human tribal society in which the behaviour of each member is constrained by internalized norms that are reproduced in each member (in the case of internal distributed management, the behaviour of every agent in the organization is controlled and coordinated by a system of constraints that is reproduced within each and every agent. As such, the constraints reach across the entire organization, and also capture the benefits [and harms] produced by their impacts on the organization as a whole). Distributed internal management can be as effective at controlling an organization as external control. But where it operates, it is often mistaken for an absence of control.

Stewart (1997, 2000, 2014) examines in some detail how the coincidence of interests between management and the organization as a whole drive the self-organization and emergence of management (management can appropriate greater resources from an organization that is managed in ways that overcome the cooperation barrier).

From the broader perspective developed here, the huge literature on the emergence of cooperation can be seen as a search for situations where constraints just happen to exist that allow some degree of consequence-capture. In contrast, 'nature' has not left the existence of suitable constraints (and the emergence of cooperation) to happenstance. Evolvable management enables the discovery and implementation of whatever sets of constraints will maximize appropriate consequence-capture in any organization in any situation.

### **Implications for Growing Artificial Life Systems**

**The origin of life.** A thorough understanding of how life originated and what distinguishes it from non-life is critically important to the 'artificial life' project. As many have noted, to be of greatest use this understanding needs to be 'substrate independent'—i.e. as far as possible it must identify the relationships between agents that are necessary to constitute life, rather than focus on the nature of specific agents that constitute particular instances of life.

The framework developed here can make a significant contribution to this understanding. Collectively autocatalytic organizations of molecular species can be self-producing and can evolve to some limited extent (Kauffman, 1993; Vasas, et al. 2012 and Nighe, et al. 2015). But the cooperation barrier seriously limits the extent of the possibility space that these organizations can explore. As we have seen, the barrier can be overcome by the emergence of appropriate management. Prime candidates for the emergence of management are coalitions of RNA molecular species that are collectively autocatalytic (Higgs and Lehman, 2015). Initially, RNA coalitions might simply have plundered the contents of other collectively autocatalytic organizations, using them to assist their own reproduction and then moving on to plunder other organizations. Importantly, these RNA coalitions would not have participated in the catalytic interactions and relationships that occurred within the organizations they exploited. They would have stood outside them dynamically and appropriated the resources they needed. The capacity to do this would have given them the *potential* to control and manage an organization as a proto-metabolism.

How might these RNA coalitions actually become managers? What would drive the transition? It is conceivable that coalitions could achieve an advantage if they discovered ways to use their catalytic capacities to enhance the productivity of an organization and manage it as a protometabolism. The existence of the cooperation barrier provided an enormous potential for RNA coalitions to do this. RNA coalitions could discover ways to intervene in organizations to support molecular species that contribute to the productivity of the organization, but would not be supported otherwise. And they could degrade or otherwise suppress free-riders that impede productivity. As a result of selection favouring RNA coalitions with enhanced management capabilities, coalitions could increasingly move away from plundering and destroying organizations. Coalitions could evolve increasingly towards a situation in which each coalition managed a particular organization as a proto-metabolism, thereby enhancing the productivity of the organization and increasing the resources that the coalition could harvest on an on-going basis. A coincidence of interests would arise between the coalition and the proto-metabolism it manages. This evolutionary sequence is broadly analogous to the historical transition which was undergone by Mongol tribes: they began as plunderers that destroyed other societies and then moved on to new conquests and pillaging. But eventually the Mongols became rulers of the societies they conquered, introducing systems of governance (management) that enhanced the productivity of the societies. Rather than plunder a society once, they could harvest an enhanced stream of benefits from it on an on-going basis.

The transition from chemistry to life. Effective, evolvable management (whether RNA or otherwise) would have enabled self-producing organizations to transition from non-living chemistry to life. As we have seen, un-managed, selfproducing chemical organizations are only able to explore a possibility space that is seriously limited. But effective, evolvable management changes everything. It opens up enormous new areas of possibility space to self-producing organizations, enabling them to go far beyond what is possible through un-managed chemical interactions and processes. Management opens the door to entirely novel and hitherto unknown arrangements of matter that are self-producing. It does so by controlling and manipulating chemical processes so that they serve the organization's functions and purposes. The nature and functioning of the constituents of the organization are no longer determined by chemistry alone. It is now dictated by the evolutionary needs of the organization as a whole. With comprehensive consequence-capture, the constituents of selfproducing organizations will tend to adapt in ways that serve the interests of the organization. As a consequence, managed organizations will tend to evolve and adapt as coherent wholes that can develop all the characteristics of individuality. In contrast, un-managed autocatalytic organizations are like ecosystems-they contain autocatalytic cycles and processes but do not evolve as individuals (comprehensive management and consequence-capture are prerequisites for the full emergence of individuality). In the service of their individuality, managed organizations would explore an extensive new space of possible organizational forms, relationships, processes and subsystems. These could not arise through normal chemical processes in the absence of

management. Management is the key to the transition from non-life to life (Stewart, 1995, 1997, 2000 and 2014).

From this perspective, the central function of DNA (and RNA before it) is not the storage of information. Its primary significance in the evolution of life is to serve as management that enables the cooperation barrier that separates chemistry from life to be overcome. The storage of information is incidental to the primary function of DNA/RNA which is to manage. Effective management requires memory.

It is also worth noting that the 'Managed Metabolism' hypothesis for the origin of life that I have presented here differs markedly from the standard version of the 'RNA World' hypothesis (Gilbert, 1986). The standard RNA World hypothesis is that life began with the emergence of selfreplicating RNA molecules that were naked (i.e. without associated metabolisms) and that then proceeded to progressively create around themselves a complex, supporting metabolism. In contrast, the Managed Metabolism hypothesis is that RNA molecules (or other potential managers) arose with and eventually took over and managed pre-existing, selfproducing chemical organizations that became protometabolisms. The Managed Metabolism hypothesis is significantly more plausible because (1) collectively autocatalytic chemical organizations are likely to have selforganized readily in organic-rich soups (e.g. Kauffman, 1993) [and their existence is likely to have been essential for the emergence of RNA self-replicators in the first place]; and (2) potential managers are much more likely to have taken over and managed these pre-existing organizations than to have created them afresh (particularly given the difficulties of building highly complex, dynamical organizations from scratch using an evolutionary mechanism that operates 'top down' and generally makes only one small change at a time).

In general, the designers of artificial life systems tend to have sidestepped the cooperation barrier that applies to the origin of life. They have done this by starting off their systems with artificial agents that can adapt as individuals without any internal restrictions on the novelty that they can produce. However, if a goal of designers of artificial life systems is to emulate the open-ended evolution of natural systems, this only postpones the necessity to deal with cooperation barriers. It merely exports the cooperation barrier to the next level-i.e. a cooperation barrier will be encountered when cooperative organization begins to emerge amongst the artificial agents that comprise the system. If artificial life systems are to be able to overcome this and subsequent cooperation barriers, they will need to be designed to enable the emergence of management along the lines outlined in this abstract. This is equally the case for any artificial life systems which seek to emulate the fundamental transition from nonlife to life.

#### References

Bagley, R. J. and Farmer, J. D. (1991). Spontaneous Emergence of a Metabolism. In Langton, C., Taylor, J., and Rasmussen, S. editors, *Artificial Life II*, pages 93-141. Addison and Wesley, New York.

Gilbert, W. (1986) Origin of Life: The RNA World. *Nature*, 319:618. Higgs, P. G. and Lehman, N. (2015). The RNA World: molecular

cooperation at the origins of life. *Nature Reviews Genetics*. 16:7-17. Kauffman, S. A. (1993). *The Origins of Order: self-organisation and selection in evolution*. Oxford University Press, New York. Maynard Smith, J. (1979). Hypercycles and the origin of life. *Nature*, 280:445-446.

- Maynard Smith, J. and Szathmáry, E. (1995). *The Major Transitions in Evolution*. Freeman, Oxford.
- Nighe, P., Hordijk, W., Kauffman, S. A., Walker, S. I., Schmidt, F. J., Kemble, H. A., Yeates, J. A., and Lehman, N. (2015). Prebiotic network evolution: six key parameters. *Mol. Biosyst.*, 11:3206-17.
- Salthe, S. (1985). Evolving Hierarchical Systems. Columbia University Press, Columbia.
- Stewart, J. E. (1995). Metaevolution. J. Soc. Evol. Syst., 18:113-114.
- Stewart, J. E. (1997). Evolutionary transitions and artificial life. Artificial Life, 3:101-120.
- Stewart, J. E. (2000). Evolution's Arrow: the direction of evolution and the future of humanity. Chapman Press, Canberra.
- Stewart, J. E. (2014). The direction of evolution: the rise of cooperative organization. *BioSystems*, 123:27-36.
- Stewart, J. E. (2015). The self-organizing society: a grower's guide. Available at SSRN: <u>http://ssrn.com/abstract=2657948</u>
- Taylor, T. (2015). Requirements for open-ended evolution in natural and artificial systems. Presented at EvoEvo Workshop at the 13<sup>th</sup> European Conference on Artificial Life (ECAL 2015), University of York, UK. Available at: <u>http://arxiv.org/abs/1507.07403</u>
- Vasas, V., Fernando. C., Santos. M., Kauffman, S. A. and Szathmáry, E. (2012). Evolution before genes. *Biology Direct* 7:1.