

Continuously evolving rewards in an open-ended environment

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Abstract

Unambiguous identification of the rewards driving behaviours of entities operating in complex open-ended real-world environments is difficult, in part because goals and associated behaviours emerge endogenously and are dynamically updated as environments change. Reproducing such dynamics in models would be useful in many domains, particularly where fixed reward functions limit the adaptive capabilities of agents. Simulation experiments described here assess a candidate algorithm for the dynamic updating of the reward function, RULE: Reward Updating through Learning and Expectation. The approach is tested in a simplified ecosystem-like setting where experiments challenge entities' survival, calling for significant behavioural change. The population of entities successfully demonstrate the abandonment of an initially rewarded but ultimately detrimental behaviour, amplification of beneficial behaviour, and appropriate responses to novel items added to their environment. These adjustments happen through endogenous modification of the entities' reward function, during continuous learning, without external intervention.

Keywords: Reinforcement Learning, Reward updating, Evolution, Open world

1. Introduction and context

This paper addresses the difficulty of simulating agents which operate in complex, open-ended environments, where the goals and objective functions of the individuals are incompletely defined. Real-world entities in open environments (e.g. humans, animals) may face multiple concurrent challenges and undertake a broad set of potentially competing tasks. They must learn how to behave without the opportunity of replaying and improving upon past choices, operating on a *linear timeline*. Changes in the environment, and the co-evolution of interactions between multiple individuals, mean the behaviour required for survival may change significantly over time. In the case of biological life, evolution provides opportunities for long-term adaptation, while short-term learning drives behavioural responses, potentially over multiple lifetimes through 'cultural inheritance'. This paper is an attempt to capture this rather constrained adaptability within a simulated environment, using embedded reinforcement learning (RL) (Arulkumaran et al., 2017) agents and a simplified evolutionary process. In RL, scalar-valued rewards serve as feedback to guide learning, typically generated externally to the agent using human-specified reward functions. Achieving the desired behaviour reduces largely to the problem of defining appropriate reward functions, which ideally provide effective (frequent and unambiguous) training signals. In well-defined tasks with clear goals ('achieve the highest score in game X', or 'place object A at position B') shaped external reward functions designed by humans can be highly successful, for example Mnih

et al. (2015); Gu et al. (2016); Silver et al. (2016); Jumper et al. (2021). Taking this further, when the goal of agents can be explicitly stated, proximity of learnt agent behaviour to the desired goal can provide a signal with which to optimise reward functions, further improving performance (Faust et al. 2019; see also the *Problem statement* below). However, in many complex open-ended environments it is not possible to know the reward functions of relevant entities. An alternative approach to designing or optimizing reward functions is to infer an agent’s objectives, values, or rewards from observation of example behaviour, using *Inverse reinforcement learning* (Arora and Doshi, 2021). While powerful in many contexts, this approach has a number of significant difficulties in the present case: it cannot produce reward functions relevant to novel (unseen) situations and faces challenges when example behaviour is ambiguous or suboptimal; it struggles computationally in high-dimensional spaces and can misinterpret actions driven by unseen features; the method can yield multiple plausible reward functions, potentially over-fitting to limited data (Arora and Doshi, 2021; Dexter et al., 2021). In the present open-ended context, *a priori* observations are not assumed available, and are of course not possible for novel environmental contexts. If a relevant dense reward function is not known, and can not be estimated, a possible solution is to embody high level goals as the reward. For example in the case of a simulated animal, a large reward for each surviving offspring might encourage multiple required subsidiary behaviours to be learnt. A drawback is the sparse reward signal coming from the environment: multiple (un-rewarded) behaviours are needed, potentially multiple times, before the high level reward can be obtained and it is unlikely these would be undertaken by chance, hence slowing the learning process. More fundamentally, the problem of how to choose the high-level reward to drive the lower level behaviours persists. The problem of sparseness can in some cases be addressed through *internal/intrinsic* rewards, which encourage exploration of novel behaviours and system states (Lopes et al., 2012; Poupart et al., 2006; Campero et al., 2020; Karayanni and Nelken, 2022) and/or improvements in knowledge of the environment (Schmidhuber, 1991; Chentanez et al., 2004; Stadie et al., 2015; Houthoofd et al., 2016). Curiosity-driven exploration is one such approach (Oudeyer et al., 2007; Oudeyer and Kaplan, 2007; Pathak et al., 2017; Zheng et al., 2021), where rewards reflect how well the agent predicts the outcome of its actions or how surprised it is by its observations. Over repeated periods of learning these approaches can yield impressive results, as in Pathak et al. (2017). In ‘novelty search’ (Risi et al., 2009; Lehman and Stanley, 2011; Conti et al., 2018), rewards are earned by diverging from stored memories of prior behaviour, and the novelty metric creates a constant pressure to innovate through state exploration. In the present case, an expected drawback of these approaches is they are not guaranteed to consistently find and exploit specific objectives, particularly for more complex problems. Behaviours which are ‘good’ in some sense for agents (for example, in aiding survival) may be abandoned for more novel behaviours. At a deeper level, it remains the case that prior decisions are made which to a large extent direct the behaviour of the agents, here deciding time should be spent exploring above all else, which may not reflect the behaviour of real-world agents in linear timelines. Additionally, human-chosen extrinsic rewards may still be required in order to achieve specific goals (such as producing offspring) as a drive to explore will alone not necessarily lead to discovery of behaviours unrelated to the agent’s own ability to explore. In the realm of evolutionary algorithms, rewards directly inspired by the survival challenges of living entities can be operationalized through an explicit (modeller-defined) fitness function

which promotes selection for progressively fitter generations, as for example in Moriarty et al. (1999); Lambora et al. (2019); Faust et al. (2019); Hallawa et al. (2020). In the present work importance is given to avoiding the modeller setting permanent objectives (defining *success*), as this would ultimately be responsible for directing behaviour, and therefore not necessarily adaptive to novel environmental conditions. An alternative to this is to use survival itself to implicitly define success (only those which behave ‘well’ will survive). This essentially calls for adaptive agents which determine their own goals, transcribe these in to dense external rewards, and promote the learning of beneficial behaviours on a linear timescale. A complete approach would also generalise beyond adjusting rewards for existing behaviours, to including means of discovering and rewarding new actions in appropriate ways.

This contribution Through simulation, this paper shows endogenous updating of a dense external reward function can produce simulated populations of agents which dynamically adjust to challenges within an open-ended environment: a self-replenishing population of agents continuously learn and dynamically adapt their rewards (and therefore their behaviour) as time proceeds. This is in contrast to existing RL approaches that attempt to find optimal policies for fixed reward functions. While agent behaviour can adapt to variations in its environment during standard RL training (and potentially in subsequent inference), because the reward function remains fixed, behavioural adaptation is essentially finding different solutions to meet the same objective, rather than reformulating the objectives (through the reward function) to meet the requirements of new challenges. An abstract ecosystem-like simulated environment is used to test these ideas: entities with finite lifetimes possess initial reward functions comprising multiple components; entities inherit (from their maternal parent) the value for each reward coefficient and also an expectation over time of rewards for each component; when they become parents they pass on an updated version of both the reward coefficient and its expectation; the modification is based on their own experience. This process is termed RULE: Reward Updating through Learning and Expectation (Bailey, 2023). This paper presents a conceptually simple test of this approach in which detrimental/advantageous options are made available to simulated entities, who adjust their reward functions in ways which are advantageous to them in avoiding extinction, without being explicitly encouraged to survive.

1.1 Scope and Problem statement

The context of this work is the behaviour of entities, and populations of entities, operating in open world contexts, with constraints on, for example, information availability, interaction rates, energy use/availability, and transfer of currencies (in the broad sense), which affect the ability of entities to survive. The environment is open in the sense there is no clear task to complete, and there are multiple simultaneous influences on entities, with the possibility of new phenomena at any time. These features are seen in many real-world contexts including ecological-, economic-, and social-systems and the generalisability of the current findings is discussed in later sections. An abstract ecosystem-like model is used here as a test case, which hosts a single type/species of agent entity (named ‘Ents’) and a single type of primary producer (‘PP’).

Problem statement The true state of the world is only partially observable by each Ent agent in the simulation. The learning problem conforms to a Partially-Observable Markov Decision Process, described by the tuple $\langle S, A, O, D, R, \gamma \rangle$, where $S = \{s_1, s_2, \dots, s_n\}$ is a set of partially observable states, $A = \{a_1, a_2, \dots, a_m\}$ is a set of actions, and $O = \{o_1, o_2, \dots, o_k\}$ is a set of observations. $D : S \times A \times S \rightarrow [0, 1]$ models the unknown system dynamics as a transition distribution. The immediate reward, $R(\theta, s, a) : S \times A \rightarrow \mathbb{R}$, where $\theta \in \Theta$ are the reward coefficients, is then discounted at a rate $\gamma \in [0, 1]$. RL is an attempt to find a policy π_R^* that maximizes the expected cumulative discounted reward R of trajectories \mathcal{T} drawn from a set of initial conditions $N \subset O$, such that $\pi_R^* = \arg \max_{\pi} \mathbb{E}_{\mathcal{T}(N, D, \pi)} \left[\sum_{i=0}^{|\mathcal{T}|} \gamma^i R(\theta, s_i, a_i) \right]$. The behaviour of the agent is therefore dependent on the reward function R , which is typically shaped through human selection of θ to result in the desired outcomes. The success of any policy can be quantified as $\mathcal{J}(\pi) = \mathbb{E}_{\mathcal{T}(N, D, \pi)} [\mathcal{G}(\mathcal{T})]$, where \mathcal{G} is the metric for trajectory success. In unambiguous RL tasks, the relationship between R and candidate (human-created) evaluation functions \mathcal{G} can be straightforward, and the chosen \mathcal{G} can even be used as a fitness metric against which to evolve successful reward functions (Faust et al., 2019). Defining success for an agent (or population of agents) in real-world open-ended systems is not so straightforward, as multiple related tasks are underway simultaneously and both the properties of agents and the challenges they face may change over time. Further, the agent may face both behavioural and evolutionary trade-offs which are not explicitly known, and maximizing lower-level rewards may have impacts on achieving higher-level success. For example, maximizing lower-level rewards for consumption might produce longer-lived entities and therefore yield larger more successful populations (higher-level success) even though population size/persistence is not part of the reward definition. Such high-level success could in principle be measured by a suitable metric, but finding a unique such function (\tilde{G}) is not expected to be possible in real-world open-ended systems (e.g. finding a unique goal and therefore suitable success metric for an animal within an ecosystem). The present work is not therefore an attempt to find an optimally-shaped reward function (i.e. $\arg \max_{\theta} \mathcal{J}(\hat{\pi})$) that maximizes an explicit success metric. Instead, the goal is to produce an emergent process which endogenously updates θ in order to maximise the unobservable success metric (\tilde{G}). Ideally, the process must explore and adjust both the reward coefficients (θ) and the basic properties of the agents (P) to prevailing conditions. Given that high level objectives emerge dynamically from contingent (survival) challenges faced by populations of entities, it is not guaranteed that endogenous processes (through selection pressures) acting on θ will result in convergence to optimal solutions, $\tilde{\pi} = \arg \max_{\theta} \tilde{\mathcal{J}}(\tilde{\pi})$, where $\tilde{\mathcal{J}}(\tilde{\pi}) = \mathbb{E}_{\mathcal{T}(N, D, \tilde{\pi})} [\tilde{\mathcal{G}}(\mathcal{T})]$. Indeed, in real-world complex adaptive systems there is ample evidence of entities occupying local physiological and behavioural minima, for example due to inescapable physiological inheritance (Wedel, 2011) or behavioural path-dependence (Heino et al., 2023).

2. Simulated open-world environment

In the simulated environment, discrete individual entities (‘Ents’) with finite lifetimes operate within a spatially extensive physically-based world. The environment enforces certain constraints, including physical constraints on movement, information transfer, and conservation of energy. An external source provides energy in the form of light which

powers the system, and reproducing ‘primary producers’ directly use this light to create biomass. All actions have an energy cost and are therefore limited by energy availability. Mobile entities (‘Ents’) with finite lifetimes exist in this environment, obtain energy through consumption, and have the potential to reproduce. They possess multiple senses and (using reinforcement-learning, RL) learn to map their observations to a fixed set of available actions which include movement, manipulation of objects, emission of signals, and the transferring of various currencies to other Ents. Some of the state variables describing each entity are fixed throughout each lifetime and inherited during reproduction through ‘genes’ from the two required parent Ents. The demands of the environment naturally produce selection pressures, and these result in evolution. The following sections provide brief descriptions of the components of the model. Additional details are given in Appendix A.

Simulated physical environment The simulated environment is a continuous 3D space, with a basal surface area of $20 \times 20 \text{ m}^2$ and unbounded in the vertical dimension. Time proceeds discretely in integer steps $\Delta t = 0.02 \text{ s}$. Basic physics (gravity, inertia, response to force, drag and scattering during collisions) is applied to all rigid objects (the simulation was built using the Unity 3D game engine, with physics updates implemented using Nvidia PhysX). Time is defined in seconds, mass in kg, and length in m. Energy is normalised to the power of the external source, S , which remains fixed at 1 throughout all of the reported experiments. The main physical elements of the world are shown in Figure 1.

2.1 Primary producers (PP)

Primary producers are the base of the food chain and produce biomass (bodymass and fruits) using energy derived directly from the external energy (light) source. Information contained in their genes entirely determines their physiology and behavioural responses. PPs grow from seeds ejected within fruits by a parent PP. On the ground, fruits notionally decompose and their seeds lay dormant for a genetically defined period before starting to grow. While on the ground, some fraction of the non-palatable fruits are available and may be collected by Ents (and potentially converted later to *processed food*). Once a seed begins to grow, it grows at a rate proportional to its mass and the strength of the incident light, until it reaches a genetically-set maximum mass. While alive, PPs compete with each other for space, within a competition radius proportional to their height, with the larger of the two killing the smaller. PPs can also be consumed directly by Ents, who remove bite-sized amounts of mass. The palatability of primary producers to Ents (and hence the sign and proportion of the biomass-derived energy available to the Ent) is determined by the genes of both Ent and PP. Eventually, PPs die for one of three reasons: (i) reaching their genetically-determined maximum lifespan; (ii) losing all mass to consumption; or (iii) being outcompeted for space by a larger nearby PP.

2.2 Ents

Simulated entities (‘Ents’) inhabit the surface of the environment and are essentially spherical in shape and have a density equivalent to that of water. To aid visual interpretation, additional body parts (eyes, crests, and tails) are given to Ents but these serve no functional purpose. The internal physiological workings of Ents are not resolved, and the ability to

perform essential physical operations, such as movement, digestion, growth and reproduction, are assumed. Ents can be damaged through impact, predation or mass loss due to starvation and can therefore die before their maximum potential lifetime is reached. Ent behaviour is learnt collectively using reinforcement learning (details below).

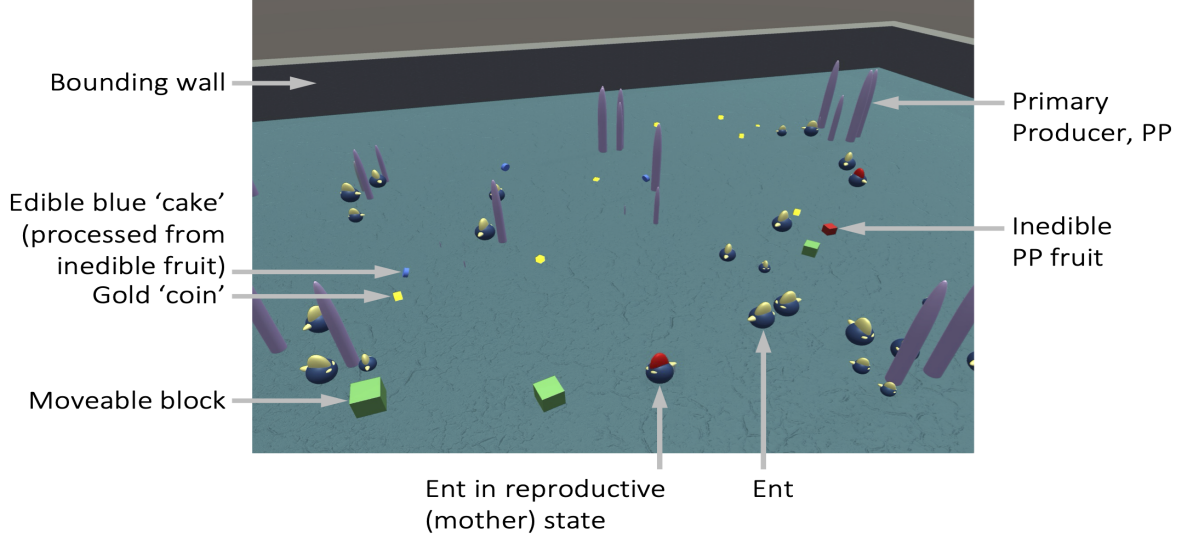


Figure 1: A screen capture of the model running, showing the environment components.

Energy Energy costs are associated with various activities, including the base running cost (resting metabolic rate, proportional to $\text{mass}^{2/3}$), movement (proportional to mass), growth of biomass, synthesizing of currencies, processing raw material (from inedible fruits to processed blue ‘cakes’), currency transactions, and collision damage (which is proportional to the strength of the impact above a minimum threshold). Ents obtain energy through the consumption of primary producers (PPs), other Ents, and processed ‘cakes’. To consume food, Ents must physically touch the object and *bite-off* an amount of mass in proportion to their own body mass. This ingested material is then notionally stored within the Ent’s stomach and gradually digested to obtain energy. The energy release occurs incrementally over a period that is proportional to the mass consumed. Further consumption is only possible once the stomach has emptied sufficiently. The energy obtained from consumption depends on the energy density of the material and its palatability to the Ent. Palatability is determined by the genetic makeup of both the Ent and material consumed (the palatability of a specific potential foodstuff is therefore subject to evolutionary adaptation); three values within the genes of Ents and PPs determine their *tissue* makeup, and are also translated to RGB to render the Ent/PP colours in visualisations. Energy obtained by digesting consumed material can be stored (as ‘fat mass’) for later use in powering physiological functions, and meeting other costs, or it can go to growing biomass (both for the Ent and any offspring). If an Ent is not fully grown, and its stored energy level is below a genetically-set amount, energy from digestion is partitioned between growth and storage according to a genetically determined ratio; if the energy store has reached the minimum amount (and the Ent is not

fully grown), all energy is used for biomass growth; finally, if the Ent is fully grown, all energy sent to the store for later use. In the absence of available food, Ents automatically draw energy from their energy stores. If no food is available and the energy (fat) store is depleted, the Ent’s remaining (lean) body mass is internally converted into energy until it reaches a minimum viable body mass (relative to its maximum previously-achieved mass). Once the minimum viable body mass is reached, the Ent dies.

Currencies By definition, interactions between entities in real systems involve a transfer of some combination of information, material and/or service (loosely termed), and to augment potentially meaningful interactions between Ents, a generalised representation of currencies is introduced in to the simulation. Here, a currency is something that one entity can choose to give to another, and is not necessarily tangible. Two key properties are used to differentiate four types of currencies: ‘subtractability’ (*True*: the currency is used-up when given; *False*: the currency is not used-up when given) and ‘synthesizability’ (*True*: currencies can and must first be created by individuals, with some material/energy cost; *False*: the currency can only be exchanged once collected). Exchange of currencies may incur a transaction cost, and rewards are associated with their transfer. One of each currency type is used in the current experiments, summarised in Table 1.

Currency	Subtractable	Synthesizable	Model component
C_1	True	True	An item requiring energy + raw material to produce. Here, inedible fruits from PPs can be collected and processed by Ents, producing processed material (‘cakes’) with maximum possible nutritional value, as currency C_1
C_2	False	False	A service-/action-related currency, which is innate and entails no additional cost (grooming, and the display of positive emotions, are examples).
C_3	False	True	A non-innate skill which can be obtained at some production cost, but is then infinitely reusable (a skill such as the ability to play music is an example).
C_4	True	False	Found objects, typically finite. Here, ‘coins’ are dropped in to the environment periodically, and can be picked-up, dropped again, or given to other Ents; uncollected coins vanish after a set duration.

Table 1: Properties of transferable currencies

Life-cycle, reproduction and inheritance Each Ent has a finite lifetime, during which it can be both a ‘father’ and ‘mother’ to offspring. Reproduction is not a specific single action in its own right, rather reproduction happens as two Ents touch when either/both of them are in a reproductive (‘mother’) state. Each Ent possesses an inherited set of numerical

values that determine its core properties, referred to as its gene, $G = \{g_i, \dots g_{n_g}\}$ (noting this gross simplification compared to biological genes). The reproductive (‘mother’) state can occur up to a genetically-determined number of times over each lifetime if/when the Ent reaches a minimum relative size and accumulates sufficient stored energy. Ents can (and do) learn to recognise when others are in this reproductive state. If ‘Ent-A’ (*father*) touches a reproductive ‘Ent-B’ (*mother*), there is an exchange of genetic material ($A \rightarrow B$) and ‘Ent-B’ produces a genetically-mixed offspring (and burdens the associated energy cost of biomass production) (genetic noise is also added during reproduction). The ‘decision’ to mate is therefore expressed through movement. Offspring are born immediately on mating and possess an energy reserve (genetically set) donated by the mother sufficient under initial gene values to keep them alive for ~ 10 s ($\sim 5\%$ of their maximum lifetime) without additional consumption. Offspring possess no other currencies or items, and these must be acquired during their lives. Ents finally die due to either (i) reaching their (genetically-determined) maximum lifespan, or (ii) loss of mass due to predation or starvation; their biomass is removed instantly from the system, along with their stored currencies and any collected items (with the exception of coins, currency C_4) with persist for 1 s before disappearing).

3. Agent behaviour

3.1 Reinforcement Learning

The ability of Ents to learn (continuously) is an important feature of the present work, and Ent behaviour (the choice of action taken) is learnt from experience using deep RL, as outlined in the introductory section. In some of the experiments described below, learning is continuous meaning decisions are part of the learning process and any change in reward function has potential to feed through to changes in behaviour. In other experiments, a previously learnt policy is used (in inference mode) to decide actions, without being updated. A high level summary of the main component of the process are provided below (technical details of the model and RL process are given in the Appendix).

Observations Each 0.02 s time step, entities make a range of observations covering their own internal state and their detectable environment (including other entities). **‘Movement’**: horizontal movement (x and z velocity) and rotation angle; **‘Vision’**: Ray sensors provide information about visible object type, direction relative to the forward direction of the Ent, and distance (17 rays, equally spaced over ± 40 degrees of the forward direction, with a range dependent upon the illumination level, up to a maximum range of 50 m). **‘Taste’ / palatability**: when Ents consume material, they sense its palatability (based on the paired sums of the three ‘tissue’ genes described above). **‘Touch’**: Ents can sense the impact damage when they hit any object through an energy cost which is proportional to the impact force (notionally associated to the repair cost). **‘Hearing’**: Within a genetically-set radius, Ents can identify signals (equal to $\{0, 1, 2\}$), and perceive their direction and distance ($s = 0$ is also the sound observation made in the absence of any Ent signals). No intrinsic meaning is given to these signals and Ents are free to use them, or not, and to make their own associations. **‘Smell/scent’**: Ents can detect the ‘scent’ of the closest Ent within their sensing radius. The ‘scent’ is a single value, calculated as the mean absolute difference between each of three ‘tissue genes’ described above. **‘Nearby group size’**: the (normalized)

number of Ents within sensing radius; **‘Inventories’**: the amount of each Currency stored (including the amount of stored energy, C_0), the ‘starvation level’ (proximity to minimum viable bodymass), amount of stored raw material (PP fruits) and processed material (‘blue cakes’), and numbers of cubes held (0 or 1). **Physiological state**: whether ready to mate and give birth (indicated by a red crest); whether ‘feeling pain’ (from being eaten, or from dying before the genetically-determined maximum lifespan); **Light level**: the intensity of the external light source.

Actions The ‘physiological’ processes of Ents (digestion, energy partitioning, growth, gestation) are regularly updated automatically without any direct control by the Ent. Ents can however choose to interact with the environment through various discrete actions (each of which entails an energy cost): moving, picking-up/dropping objects, exchanging currencies and signals, eating, processing raw materials and reproducing. Any set of non-exclusionary actions can be taken in each successive 0.02 s time increment, as listed here (details in Appendix A). Those actions which are not logically possible are ‘masked’ (made unavailable) to reduce the action space for learning. For example, the action to drop an object is masked if the object is not currently held. **Move**: Ents can choose to rotate in 1° increments around their vertical axis, $\Delta Rotation = \{0, 1^\circ, -1^\circ\}$; a forward force $\Delta F = \{0, \delta F, -\delta F\}$ can be applied, where the increment δF is a function of Ent lean mass, while the force acts on the total body mass. **Eat/bite**: $\{true, false\}$ if Ents are touching a ‘soft’ object they can bite it and take some of its mass. If the palatability score is positive, the material is digested over a period of time (during which no further consumption is possible) and energy is obtained (depending on the energy density and palatability of the material); if palatability is negative, no energy is obtained from digestions and a ‘poisoning’ energy reduction is applied, in proportion to the palatability score. **Emit signal**: Ents choose whether to remain silent (emit signal 0), or to emit signal $\{1, 2\}$. **Give currency**: When touching another Ent, any currency possessed can be given to that Ent, in increments of C_{given} , which is determined genetically (G[66-69]) for each currency (except for currency C_4 (coins) which is given one-at-a-time); the receiving Ent increases its currency inventory by this amount; the giving Ent reduces its inventory by this amount only if the currency is ‘subtractable’, and by zero if the currency is ‘non-subtractable’. **Pick up object**: When touching any movable object with mass less than the Ent, a $\{true, false\}$ choice is made to pick it up; all objects other than blocks can be stored in the Ent’s inventory; blocks are carried aloft by the Ent and in this case no other objects can be picked up until the block is dropped. **Drop object**: any object being carried (a cube, coins (C_4), fruits or processed ‘cake’ material) can be dropped, with a $\{true, false\}$ choice. **Process raw material in inventory**: If the Ent is carrying a PP seed, the mass of the seed can be converted to a processed ‘blue cake’, which is edible and has the maximum possible palatability (hence, maximum possible energy yield on consumption/digestion). Once processed, the blue-cake must be dropped to be eaten. It can be dropped and picked up to store again as required.

Training Proximal policy optimisation, PPO (Schulman et al., 2017), was used to train the Ent agents, implemented through the ML-Agents Toolkit within Unity (Juliani et al., 2020). A Python API provides a low-level Python interface for communication between the simulation and relevant PyTorch libraries. In the present case, only a single type of Ent is present in the simulations and PPO is implemented with a single policy, with all active

Ent agents contributing to the learning process and sharing that same policy (details of the chosen hyper-parameters are provided in the Appendix).

3.2 Rewards and Reward updating via RULE

In order to create a population of Ents which can potentially go on to update their reward functions endogenously, an initial reward function is required to begin the process of learning (an alternative, not chosen, being to slowly evolve such a function from scratch). Approaches to creating dynamic reward functions are discussed in Section 6. The chosen reward function (Equation 1) comprises a set of minimum essential components (for eating, reproduction and pain) plus components for each of the ($N_C = 4$) currencies, to provide potential for richer Ent interactions and experimental manipulations. The reward obtained at time step t is defined

$$R = f \cdot \theta_{\text{consumption}} + b \cdot \theta_{\text{reproduction}} + p \cdot \theta_{\text{pain}} + \sum_{j=1}^{N_C} \Delta C_j \theta_j \quad (1)$$

For each of the 7 reward components there is a coefficient θ which scales the contribution to the total. The first term accounts for consumption (of food material) where f is the food mass consumed, the second is reproduction where b is the number of offspring produced (rewards are given to both parent Ents). The third term defines the contribution of pain (due to impact damage, being bitten/eaten or dying before the maximum lifespan is reached) and is the only reward coefficient which remains fixed, providing an anchor to which all other reward coefficients are relative. The final term accounts for net changes in currencies. The proposed algorithm for endogenous reward updating is outlined in Algorithm 1 below and it results in updates to θ_i (where i indexes all components except *pain*). The algorithm can be qualitatively described as follows: each Ent inherits (from its mother) the value of each reward function coefficient θ_i and also an expectation distribution (over age) of the accumulated reward for each component. If an Ent becomes a mother, it passes on updated versions of each reward coefficient and each expectation distribution, where the modification depends on its own experience: for each reward component, if the mother Ent has accumulated more reward than expected (within a 20 s age bin), a larger reward value and increased expectation is passed on, and *vice versa*. Ents pass on reward coefficients to their offspring which encourage behaviour closer to their own than had no modifications been made. Ents which are *more successful* in some way, because of their experiences in the environment, will be more likely to pass on their encouragements, and what is effectively a majority vote on reward updates (and therefore eventually behaviour) occurs at the population level.

Algorithm 1 RULE: Reward Updating through Learning and Expectation**Require:** θ , $E(\tau)$, α , β , $\text{lifeStatus} \equiv \text{alive}$ **Ensure:** Offspring with updated Reward Expectations $E_i(t)$ and Reward coefficients θ_i , if offspring produced

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1: Initialize  $A_i \leftarrow 0, \forall i$   $\triangleright i \in [1, \text{numberOfRewards}]$ 
2: while alive do
3:    $A_i \leftarrow A_i + R_{i,t}, \forall i$   $\triangleright$  Incrementally sum rewards each time step  $t$ 
4:   if producing offspring then
5:      $\tau = f(T)$   $\triangleright$  Calculate age bin  $\tau$  based on current age of parent,  $T$ 
6:      $\Delta_i \leftarrow A_i - E_i(\tau), \forall i$   $\triangleright$  Compare accumulated rewards to expected rewards
7:     if  $\Delta_i > 0$  then
8:        $\delta_E = \alpha_i$ 
9:        $\delta_\theta = \beta_i$ 
10:    else if  $\Delta_i < 0$  then
11:       $\delta_E = -\alpha_i$ 
12:       $\delta_\theta = -\beta_i$ 
13:    else
14:       $\delta_E = 0$ 
15:       $\delta_\theta = 0$ 
16:    end if
17:     $E_i(\tau) \leftarrow E_i(\tau) + \delta_E$   $\triangleright$  Updated Reward Expectations
18:     $\theta_i \leftarrow \theta_i + \delta_\theta$   $\triangleright$  Updated Reward Coefficients
19:    Offspring. $E_i(\tau) \leftarrow \text{Parent}.E_i(\tau), \forall \{i, \tau\}$   $\triangleright$  Pass on updated values to offspring
20:    Offspring. $\theta_i \leftarrow \text{Parent}.\theta_i, \forall i$ 
21:  end if
22: end while

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Definitions: θ : the set of reward coefficients; R : the set of reward values obtained at time t ; α : the set of reward expectation increments; β : the set of reward coefficient increments; $E(\tau)$: the set of initial Expected Accumulated Rewards distributions, across age classes τ

The time complexity of the RULE algorithm depends on the number of iterations of the *while* loop (determined by Ent lifespan, of order 10e4-10e5), and per iteration on numberOfRewards, so overall $O(\text{Iterations} \cdot \text{numberOfRewards})$.

4. Experiments

4.1 Purpose and overview

To test the RULE algorithm assessments were made of whether Ent populations could make significant changes (autonomously) to their reward coefficients (and hence their behaviour, under continuous learning) in order to improve their chances of survival, in response to changing environmental conditions. In Experiments 1 & 2 outlined below, a challenge to population survival was set up which can only be overcome if the population ‘chooses’ to abandon certain behaviour, running contrary to the initial reward structure (later populations must act ‘against the will’ of the initial population). The environment of the Ents was

set up initially with constant benign conditions and initial reward coefficients (θ) hand-chosen to yield learnt behaviour conducive to a persistent (multi-generational) population. Within this setting the collection of coins (currency C_4) is rewarded. Collecting coins is an intentionally frivolous activity, providing no immediate benefit and entailing a small time/energy (and therefore opportunity) cost. During Experiment 2, coin availability was progressively increased to the point where so much time/effort could be spent on coin collection that the population would suffer (through starvation and a lack of reproduction). This is the *coins challenge* the Ent population faced. Experiment 3 is a repeat of Experiment 2 but with coin collection directly affecting Ent stored energy and reproduction probability. The purpose of this experiment is to assess the ability of Ents to adjust their behaviours due to more direct signals compared to the more implicit opportunity-cost signal associated with coin collection in Experiment 2. Where there are positive energy/reproduction benefits, coins are referred to as ‘vitamins’, and when the effects are negative, ‘poisons’. For the RULE algorithm to be judged successful, the population must survive the *coins challenge* and be able to modify behaviour in response to the opportunities and risks associated with vitamins and poisons. The purpose of the final experiment (Experiment 4) was to assess whether reward updating could ‘switch on’ behaviour associated with previously unseen actions, and is a test of the generalisability of the RULE approach.

4.2 Baseline conditions

In all model runs, both the initial Ent population size and initial PP population size were set to 100, and these were distributed randomly across the inner 7x7 m (PPs) and 6x6 m (Ents) of the 10x10 m surface of the simulated environment. PP initial age and mass were chosen at random (to avoid forcing cyclic behaviour in biomass availability): $age \sim U[1, 100]$ and $mass \sim U[5, 50]$. Ent initial ages were set to zero (so initial training is forced to cover this crucial initial period where risk of death by starvation is high), and their mass to 3.75 kg (the offspring mass of the lightest possible mother). Each Ent is provided (by the mother) with an initial energy store equal to 1 unit (this amount is genetically determined and therefore subject to change when evolution is enabled). Batches of currency C_4 (*coins*) are dropped into the environment (materializing at a random height, $\sim U[0.05, 0.1]$, and falling to the surface under simulated gravity) at regular intervals, with a batch of n_{coins} coins per 10 s (referred to as a ‘coin cycle’). Coins persist on the surface for 20 s (during which time they may be picked up) before disappearing. Under baseline conditions $n_{coins} = 10$. The external light source was held constant throughout all simulations at a value of 1. The simulation halted if the Ent population reached zero, and ran no longer than the maximum time limit of 20,000 s for all experiments.

4.3 Preliminary stage

Baseline policy training and population persistence The first step involved choosing (by trial and error) a set of initial values for genes and reward coefficients, and running RL training for a sufficient number of steps to create a robust initial policy under baseline conditions. During the training process, if all Ents in the population died, the environment was reset. Eventually, as the behaviour of the Ents improved under training, reproduction

offset the loss (death) of Ents, and the population survived for increasingly longer periods. What counts as a ‘robust’ policy was determined using two attributes:

(1) Persistence: the ability to maintain a finite population of Ents over time under benign conditions. This requires sufficient rewards for at least food consumption (for energy supply) and reproduction. The reward coefficients found to meet this requirement were $\theta_{consumption} = 0.3$ and $\theta_{reproduction} = 1$. In the same spirit, the initial genes chosen for Ents and PPs make PPs palatable to Ents. Relatively small coefficients were assigned to changes in currencies ($\theta_{1,2,3} = 0.1$), to encourage currency transfers but to ensure currency dynamics did not initially dominate eating and reproduction during the initial period. The reward for dying before the genetically-determined maximum lifespan (from a lack of mass, due either to starvation or predation) was set to $\theta_{Pain} = -1$. The reward for being bitten by another Ent was set to $\theta_{consumption} \cdot \theta_{Pain} = -0.3$. It is unlikely the initial reward coefficients and gene values are optimal in any sense and this was not the goal, but they were sufficient to produce stable populations from which the dynamic updating processes could proceed.

(2) Coin collecting: The second necessary attribute was that the Ents should expend effort collecting currency C_4 (‘coins’). Coin collection is a necessary part of the experiment outlined below and this behaviour can only be achieved if Ents are sufficiently rewarded. The reward for ΔC_4 (collection of a single coin) was set to $\theta_4 = 0.3$.

Baseline reward expectations Following training of the baseline policy (with fixed θ values) reward expectation distributions, $E_i(\tau)$, were created, as required for later application of the RULE algorithm. To achieve this, the model was run with evolution disabled (offspring received an exact copy of their mother’s genes), reward coefficients (θ) held constant, and Ent behaviour driven by the baseline policy (in inference mode, with no learning). Populations of Ents and PPs were initiated (parameter values equal to those during initial training) and all expectation values were initialised to zero. The model was run for 10,000 s, updating reward expectations for each offspring according to RULE, providing sufficient time for the expected reward distributions $E_i(\tau)$ to stabilize at consistent values (see Appendix A.2.4 for further details). These distributions, together with the trained RL policy, provide the necessary ingredients for the experiments to proceed.

4.4 Experiment design

Assessing whether Ents can survive the coins challenge involves controlling combinations of three factors: evolution (of genes), learning (continuously updating the policy through training), and reward updating with RULE (which can only happen during learning) - see Table 2. Note that the learning condition here does not involve resetting of the environment if the population dies. The population is expected to be replenished through reproduction, so if the population collapses that is the end of the experiment. Evolution can be disabled by giving offspring an exact copy of their mother’s genes (and disabling mutation); learning can be switched-off by running the Ent behaviour model in inference mode rather than training mode; finally, reward updating (even if learning is enabled) can be switched off by setting $\delta_E = \delta_\theta = 0$ for all reproductions.

Experiment 1: Baseline coins sensitivity The baseline sensitivity of the population to coin rate was measured under otherwise standard conditions, using Ents with no evolution

or learning (and no reward updating), primarily to confirm whether a ‘fatal dose’ of coins exists, and to estimate its value (to inform subsequent experiments and interpretations). The experiment involved 10 repeat runs at each of 10 different (constant) coin rates (from 0-80 coins/10 s). Each simulation was run for 10,000 s, as this was deemed sufficient time to assess with confidence whether a population could survive.

Experiment 2: Coins challenge A total of 18 combinations of conditions were used in Experiment 2. For each of the 6 conditions listed in Table 2 experiments were conducted with: (i) constant coin rate of 10/cycle (baseline control conditions); (ii) a linear increase in coin supply at a *slower* rate, where the number of coins dropped per 10 s cycle was equal to $n_{coins}(t) = 10 + 0.0097t$ (reaching 107 coins per cycle at 10,000 s); (iii) a similar linear increase in coin supply but at a *faster* rate, $n_{coins}(t) = 10 + 0.014t$ (reaching 150 coins per cycle at 10,000 s). For the RULE algorithm to be deemed successful, Ent populations must survive increases in coin supply rates which are fatal to populations under the same conditions without RULE being enabled.

Experiment condition	Evolution	Learning	Reward updating
[2a] Inference-only	False	False	False
[2b] Evolution-only	True	False	False
[2c] Learning-only	False	True	False
[2d] Evolution + Learning	True	True	False
[2e] Learning + Reward-updating	False	True	True
[2f] Learning + Reward-updating + Evolution	True	True	True

Table 2: List of experiment conditions. In each case, there are three coin rate conditions, (i) constant (10 coins / cycle), (ii) Increasing from 10 to 107 coins / cycle, (iii) Increasing from 10 to 150 coins / cycle.

Experiment 3: Vitamins and poisons The purpose of Experiment 3 was to assess the extent to which explicitly ‘beneficial’ or ‘harmful’ behaviours can be amplified/abandoned by reward updating. This is in contrast to the implicit (opportunity) cost associated with picking up otherwise neutral coins in Experiment 2. In Experiment 3, the effects of picking up coins are extended to (i) an instantaneous change in the energy stored by the Ent (ΔC_0), and (ii) a multiplier of the probability of reproduction. Under these changes, ‘coins’ are re-conceptualised as ‘Vitamins’ and ‘Poisons’: *stronger vitamins*, $\Delta C_0 = 10, P = 5$; *weaker vitamins*, $\Delta C_0 = 0.5, P = 1.25$; *stronger poison*, $\Delta C_0 = -2, P = 0.1$; *weaker poison*, $\Delta C_0 = -0.5, P = 0.5$. To pass this test, rewards for picking-up (beneficial) vitamins should increase (from the baseline value of $\theta_4 = 0.3$), and rewards for (harmful) poisons should decrease to the extent that the population is able to survive. This is a particularly difficult challenge in the case of poisons, as Ents begin the experiment with strong poison-collecting behaviour and relatively high reward coefficients for something that is highly detrimental to their survival prospects.

Experiment 4: Dormant rewards The purpose of this experiment was to assess whether (under RULE) Ents can successfully adopt/ignore beneficial/harmful activities of which they

have had no previous experience. This is in contrast to Experiment 2 and 3 which test the ability to modify already-learnt behaviours in the face of new environmental conditions. The introduced conditions chosen were the coins and vitamins of Experiment 3. Experiment 4 therefore required starting the baseline training process from the beginning to establish a persistent Ent population without coins having been present, and without coin collection having been learnt. Experiment 4 began with this persistent population, with the coin reward coefficients (θ_4) set to a low value of 0.001, and with expected distributions for coin rewards initialised randomly within the range $[-0.001, 0.001]$. In two separate experiment runs, (1) coins, and (2) vitamins, were introduced at a linearly-increasing rate ($n_{coins}(t) = 10 + 0.014t$, the higher rate used in the coins challenge). For RULE to be deemed successful in these contexts, Ents should discover and then learn to pick up (beneficial) vitamins; conversely, they should not learn to pick up detrimental quantities of (harmful) poison.

5. Results

Preliminary stage results During baseline training (with repeated periods of learning and re-setting the environment after Ent population collapse), Ents successfully learnt a policy which allowed them to survive in their environment, maintaining a persistent population through reproduction. At a minimum this required Ents to successfully identify and then move to targets, eat, and reproduce. The time series of rewards obtained during initial training is shown in Figure 2, along with data on the succession of Ent populations observed. During the early phase of training (0-30e6 steps), Ent populations collapsed after short periods as individual Ents had not learnt to eat and reproduce. As these and other skills were acquired through training, populations persisted for longer durations, producing successive generations (Figure 2b), eventually resulting in a stable population (Figure 2a) which met both the persistence and coin-collecting criteria outlined above. In subsequent experiments involving continuous learning (a requirement for RULE) the PPO algorithm (with constant learning rate) was found to be stable, with no evidence of reward/behaviour deterioration, or failure to learn.

Results for the ‘Baseline Expected Rewards’ distributions $E_i(\tau)$ are shown in Figure 3 for Currency, Consumption and Reproduction rewards. Variation was observed in the distributions of individual Ents, due to differences in life experience accumulated down the maternal line. However, over the course of the 10,000 s model run, mean $E_i(\tau)$ values (averaged over the last $n = 500$ Ent births) converged on the values shown (black squares in Figure 3) and were used as the starting distributions for each Ent in subsequent experiments. The red circles in these figures are the $E_i(\tau)$ adjustments following the ‘coins challenge’, and are discussed below.

Experiment 1: Baseline coins sensitivity Summary results for the preliminary coin-sensitivity experiment are shown in Figure 4. Part (a) plots the final Ent population (after 10,000 s) against the coin rate. The inset shows the posterior probability of population collapse. Coin rates ≤ 30 did not impact the survival probability of the Ent population; coin rates ≥ 50 always collapsed the population, and those intermediate rates produced increased risk of collapse as coin rate increased (see inset). Between model runs there was broad consistency and example population time series are shown in Figure 4(b). These results

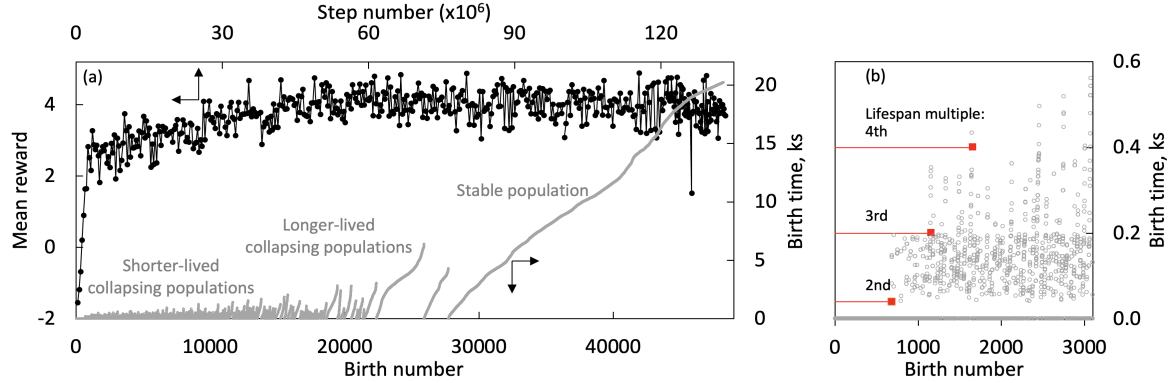


Figure 2: Baseline training. (a) Black symbols indicate the total reward obtained at each step during the initial policy training. Data in light grey are the birth number of each Ent born during the training period, plotted against ‘Birth time’ (time of birth since the beginning of the population). Stronger vertical components indicate lower reproduction rates, leading to population collapse; more diagonal components indicate more consistent birth rates. Part (b) shows the first 3,100 births in greater detail, showing where multiples of the maximum lifespan (200 s) are observed, indicating increasingly successful Ent survival and reproduction. Below “2nd”, no successful offspring are created; above “2nd” are second generation Ents; “3rd” and “4th” are not strictly generation numbers, but indicate increasing population persistence as Ents are born at times which can only be reached due to multiples of the maximum lifespan.

together demonstrate there is indeed a maximum survivable ‘dose rate’ of coins, and that increasing coins over time to beyond this limits would provide a suitable survival challenge.

Experiment 2: Coins challenge This experiment involves a linear increase in coin-rate (number of coins dropped each 10 s interval), under the conditions summarised in Table 2, Experiments[2a-f]. Figure 5 displays representative individual time series from each of these experiments, with (S) and (F) indicating whether the *slower* or *faster* increase in coin rates is shown. Table 3 provides descriptive data on the outcomes of these experiments and summary plots of repeated trials are shown in Appendix A.3. In sections A and B of Figure 5 coloured lines show data from experiment conditions (orange for populations which collapse and blue for those which survive); green horizontal lines indicate the [min,max] range of population collapse times. Grey lines plot results from control conditions where coin rates were held constant at 10 coins/cycle (noting all populations survived all control condition runs).

Section A of Figure 5 shows (in the upper row, darker orange) Ent population over time for experiment conditions 2a-2d. The lower row (lighter orange) shows the rate of coin collection across the whole Ent population, falling to zero as the populations crash. Dashed lines indicate the availability of new coins increasing over time (collection rates may exceed supply rates if coins dropped by Ents are subsequently picked-up). Results in parts (a,b) (‘Inference [2a]’, where Ent behaviour choices are made using the fixed baseline policy, with no additional learning) show the population to be initially stable as the coin rate increases linearly (collection follows supply, meaning almost all available coins are collected). Rather abruptly at ~6,000 s the population and coin collection rate fall to zero. This coincides with

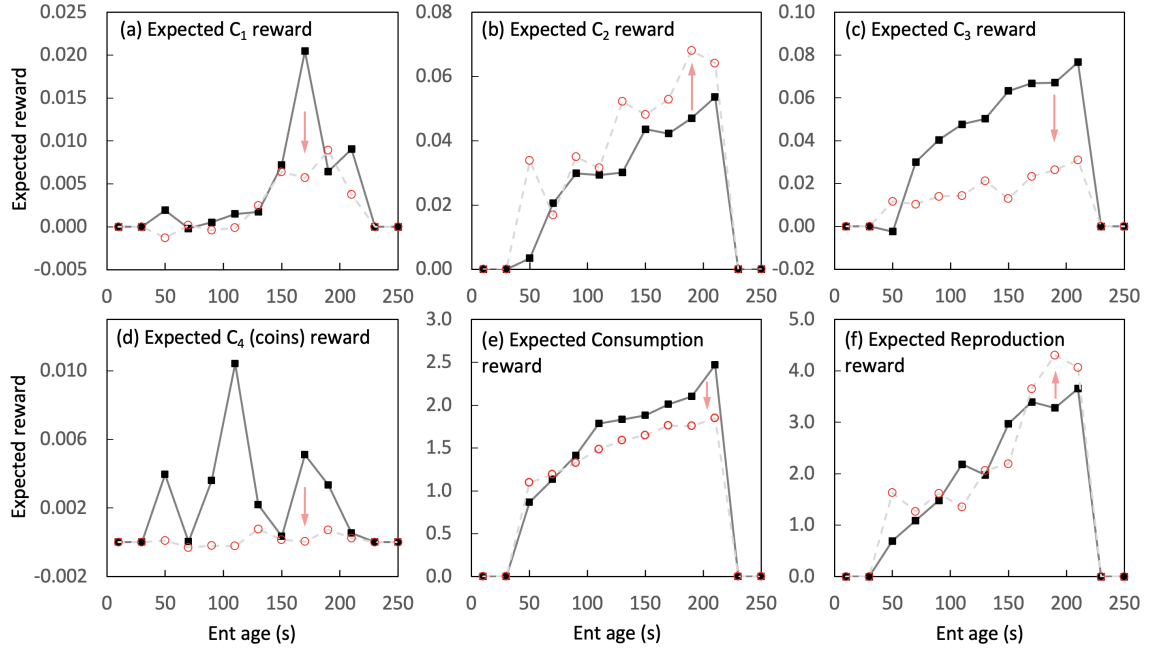


Figure 3: Expected rewards distributions $E_i(\tau)$ are shown here for Currency, Consumption and Reproduction reward coefficients. Black squares indicate the mean values (at each time point) from the last 500 Ents born in the 10,000 s simulation. These values are the ‘Baseline reward expectations’, following the initial policy training based on the chosen reward function coefficients (red circles show results from the coins challenge, discussed in the main text).

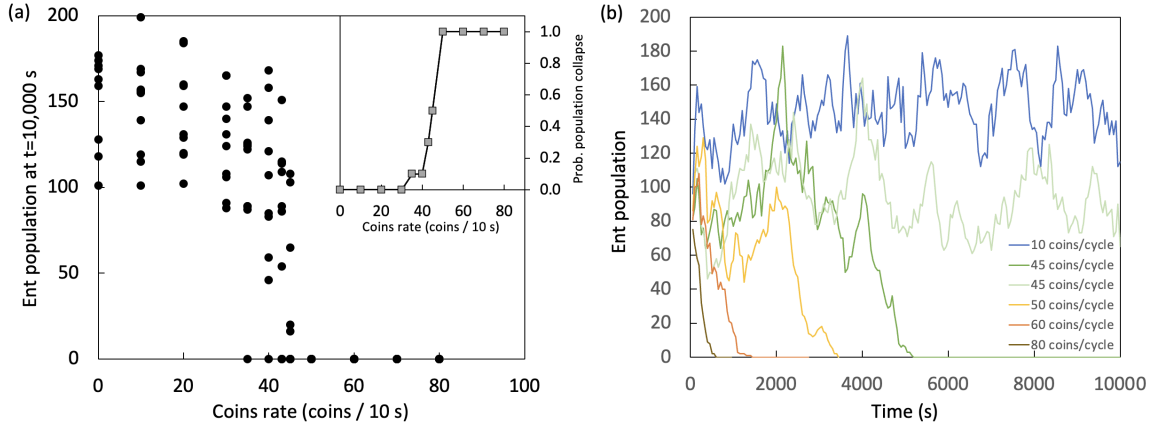


Figure 4: Results from the coins sensitivity experiment. (a) Final population after 10,000 s at the constant coin rate shown in the horizontal axis. The inset shows the probability of population collapse. (b) Example Ent population time series for a range of coin rates.

the coin supply rate increasing beyond ~ 60 coins/cycle (~ 300 coins/50 s), consistent with results shown in Figure 4. The results shown are for the slower coin increase rate but as

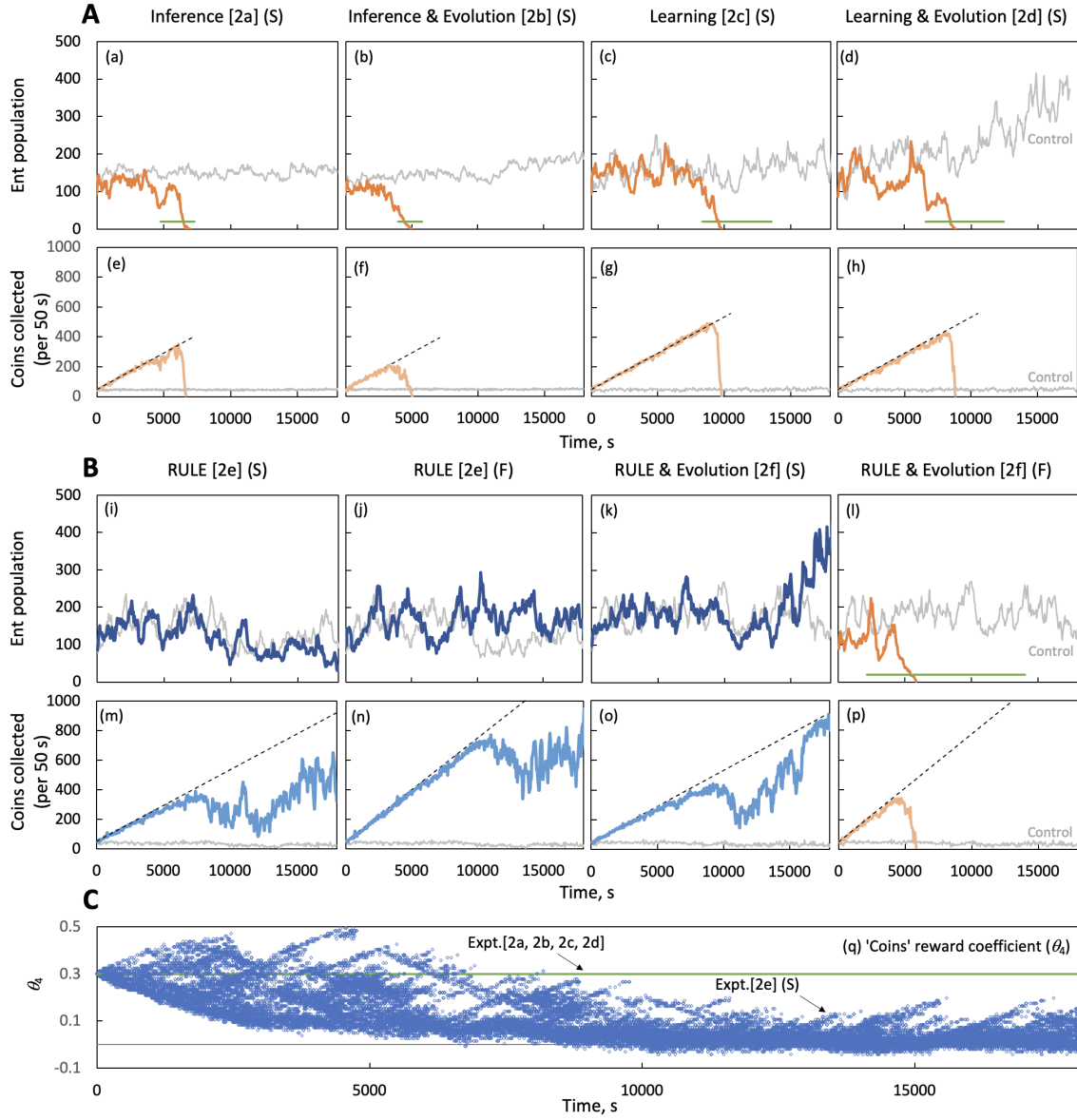


Figure 5: Results from Experiment 2: the ‘coins challenge’. Dashed black lines indicate the availability of new coins. Data are plotted as orange/blue to highlight cases where Ent populations collapsed/survived under the experimental conditions. For collapsing populations, green horizontal lines indicate the [min,max] range of collapse times from multiple repeated runs (see A.3). Grey lines are data from control conditions in which the coin rate was kept constant at 10 coins/cycle. In part (q) blue symbols indicate coin reward coefficients θ_4 for each Ent offspring, plotted against the time of its birth. Further details are available in the main text.

indicated in Table 3, ‘faster’ coin increase rates produce similar results, but with shorter-lived

populations, as the ‘fatal dose’ of coins is reached more quickly. In parts (b,f), it can be seen that enabling ‘evolution’ (Expt.[2b]) under the inference condition does not lead to longer-lived populations, and there is some evidence it may shorten survival times (Table 3). Under the ‘Learning’ condition (parts c,g of Figure 5), where Ents actively update their shared policy throughout the experiment, populations survive the coins challenge for ~50% longer (and are able to collect more coins) compared to (non-learning) ‘inference’ conditions. However, the overall pattern is similar in that populations (Figure 4c,d) and collection rates (Figure 4g,h) for both coin increase rates, with and without evolution enabled, eventually collapse over a similar range of times as coin availability increases (Table 3). Under the control conditions, there is no evidence that continuous learning was detrimental to population size (in fact the combination of Learning & Evolution led to significant population increase, Figure 4d). The important observation from Experiments [2c] and [2d] is that neither learning (with a fixed reward function), nor Learning & Evolution, provided means for the population to survive the coins challenge.

Section B shows results for Experiments 2e and 2f (which involve continuous learning with adaptive reward coefficient updating, RULE), for the faster (F) and slower (S) rates of coin increase, both with and without evolution enabled. Figures 5(i,j) show results for Experiment [2e] (RULE). Unlike results presented in part A, the large majority of populations do not collapse as coin rates increased (Table 3), with populations typically remaining at broadly consistent levels throughout the experiment. The coin collection rates initially increase, following the coin supply, but then reduce considerably, meaning not all available coins were collected. This reduction was not caused by population decline (as in the Inference and Learning conditions) but by changes in Ent behaviour driven by updating of the reward coefficient (θ_4 ; part C), associated with changes in the expected reward distributions ($E(\tau)$, Figure 3). Successful Ent populations survive increases in coin rate to levels well beyond the 50 coins/cycle limit identified in Figure 4. The majority of populations across repeat runs passed this test (see Table 3). In some cases, eventhough behavioural change occurred as θ_4 reached near-zero values, populations fell to low levels, and some collapsed. This is an expected consequence of the stochastic nature of small spatially distributed populations, where chance plays a part in, for example, the likelihood of reproduction, and is well-documented in many ecological populations (e.g. Kramer et al. (2009)). Reward updating via RULE therefore appears to provide a mechanism for populations to survive the coins challenge.

For the ‘RULE & Evolution’ condition [2f], the success of the population in defeating the coins challenge depends on the rate at which the coin supply is increased. At the lower rate (S), the response in terms of coin collection (Figure 5(o)) is similar to the ‘RULE’ condition over the first ~11,000 s, but the population increases dramatically from ~14,000 s, and the coin collection rate eventually matches the coin availability rate. In contrast, under the higher coin increase rate (F), the population collapses relatively quickly as coin availability increases (Figure 5(l,p)).

Section C of Figure 5 shows the reward coefficient for coins (θ_4) over time, for Experiments [2a,2b,2c,2d,2e]. In the case of Experiments [2a,2b] (Inference) and Experiments [2c,2d] (Learning), all reward coefficients are held constant. In Experiments [2e,2f] (RULE), where reward coefficients are updated actively over time, significant changes are observed (results for both experiments are similar (Appendix A.3) and for clarity, only example results for 2f are shown). Each point plotted represents the θ_4 value of an Ent offspring, at the time

of its birth. A dominant features is for θ_4 to reduce towards zero, and this is evident from the beginning of the time series. The range of θ_4 values initially increases, going above and below the initial value of 0.3, before becoming more restricted at later times. During the ‘coins challenge’, the increasing availability of coins means some Ents collect more, and subsequently give birth to Ents which similarly pick up relatively large numbers of coins. This drives up θ_4 for those Ents and these ‘lineages’ can be seen in the data plotted in Figure 5(i). At the same time, other Ents pick up fewer than expected coins, and give birth to Ents who have similar experiences. Hence in the early stages of Experiment 2e, there is a diversity of branching θ_4 responses. As coin availability increases further, those Ents who happen to collect more coins are on average less successful at producing offspring, and so the ‘signal’ to the population encouraging greater coin collection is relatively weak. Conversely, those collecting fewer coins will on average produce more offspring with θ and $E(\tau)$ values which encourage less coin collection. As these effects continue, θ_4 drops significantly over the course of the experiment, to values below those for both $\theta_{consumption}$ and $\theta_{reproduction}$. The outcome in terms of behaviour is that Ents pick up coins if they are in their vicinity, but won’t abandon food or reproduction opportunities for coins. So, while some coin collecting behaviour continues, many coins are left un-collected and populations survive.

Experiment		% survival (n)	$t_{collapse}$ [min, max]	θ_4 (coins)	θ_5 (consumpt.)	θ_6 (reproduct.)
[2a] Inf	(F)	0 (10)	5020 [4000,5950]	-	-	-
	(S)	0 (10)	6192 [4750,7300]	-	-	-
[2b] Inf,Ev	(F)	0 (10)	3920 [3250,5450]	-	-	-
	(S)	0 (10)	4940 [3950,5800]	-	-	-
[2c] Lrn	(F)	0 (10)	7925 [6100,10200]	-	-	-
	(S)	0 (10)	9970 [8350,13550]	-	-	-
[2d] Lrn,Ev	(F)	0 (10)	5940 [3700,8300]	-	-	-
	(S)	0 (10)	8930 [6600,12450]	-	-	-
[2e] RULE	(F)	83 (12)	-	0.04 [0.03,0.06]	0.32 [0.24,0.4]	0.97 [0.95,0.98]
	(S)	80 (10)	-	0.03 [0.03,0.06]	0.32 [0.22,0.42]	0.97 [0.97,0.98]
[2f] RULE,Ev	(F)	0 (12)	7340 [2150, 14050]	-	-	-
	(S)	75 (12)	-	0.03 [0.02,0.06]	0.38 [0.2,0.62]	0.98 [0.97,0.98]
Vitamins						
Stronger	(F)	100 (6)	-	1.0 [1.0,1.0]	0.13 [0.11,0.14]	0.99 [0.99,1.0]
Weaker	(F)	100 (6)	-	1.0 [1.0,1.0]	0.04 [0.02,0.07]	1.00 [0.99,1.0]
Poisons						
Weaker	(S)	75 (12)	-	0.01 [0.01,0.03]	0.24 [0.17,0.42]	0.97 [0.96,0.97]
Stronger	(F)	0 (12)	4988 [4400,6100]	-	-	-

Table 3: Experiments results summary. (F) and (S) refer to the slower and faster rates of coin increase (§4.4). Each experiment was repeated n times and the ‘% survival’ column indicates the proportion of those populations surviving the experimental conditions (bold font highlights those conditions for which the majority of populations survived the experiment). For conditions where all populations collapsed, the mean [min, max] time to collapse is shown. No time to collapse is shown for conditions where survival dominates. In these cases, the final θ_i values (the mean of the last 10% of each time series), and the [min,max] range are given. These data are not provided for populations which collapsed during the experiment.

Effects of evolution For experiment [2b], 10 repeats of the control condition were run (evolution was enabled, but no learning or reward updating). Across the results from these runs some gene values showed no consistent changes and drifted to both higher and lower values. In contrast, other gene values showed consistent patterns across all 10 repeat runs. The left-side (blue) panels of Figure 6 show representative results of this latter group. Each data point represents the gene value for an Ent offspring, plotted at its time of birth. The distributions shown to the right of the gene time series are frequency distributions of the gene values from the last 500 offspring of the time series; the dashed line indicates the initial genes value. Amongst these data there is a tendency over time towards the following: larger Ents [evolving from an initial value of 25 \rightarrow mean value of 36] (b) who keep larger energy reserves [1.0 \rightarrow 2.1] (f), with consistent ($<5\%$ change) relative stomach sizes [2.0 \rightarrow 2.1] (g) and maximum lifespans [200 \rightarrow 193] (c); able to reproduce when smaller (lower body mass) [0.75 \rightarrow 0.60] (d), giving birth to relatively larger offspring [0.20 \rightarrow 0.31] (e). These genetic changes yield a stable persistent population. For comparison, equivalent (orange) plots are shown on the right side of Figure 6, generated using data from Experiment [2f] (increasing coin rate, learning and reward updating via RULE & Evolution). Under these conditions, the evolutionary response has some key differences to the control condition described above: Ents become *smaller*, with *lower* maximum mass [25.0 \rightarrow 13.6] (i); their maximum age reduces to a greater extent [200 \rightarrow 118] (j); minimum reproduction mass is reduced by a much greater amount, with the distribution running in to its enforced minimum value at 0.2 [0.75 \rightarrow 0.23] (k) while the relative mass of offspring increases to a greater extent [0.20 \rightarrow 0.36] (l); relative stomach masses *increase* dramatically [0.20 \rightarrow 0.35] (n) and energy reserves (while more variable) extend to higher values [1.0 \rightarrow 2.7] (m). Proposed explanations for the changes observed during the control conditions and the coin challenge conditions are presented in the Discussion section below.

Experiment 3: Vitamins and poisons Figure 7 presents example results from Experiment 3, a ‘coins challenge’ experiment (with RULE and no evolution) in which the act of picking up coins was combined with either energy-and-reproductive benefits (coins as ‘vitamins’) or reductions (coins as ‘poisons’). The top and middle rows provide time series of reward coefficients for coins (θ_4) and food consumption ($\theta_{consumption}$) respectively; the bottom row shows Ent population time series. Three experimental conditions are shown in the columns: weak vitamins, strong vitamins, poison (see experiment description section above for details). In both ‘vitamins’ experiments, θ_4 increases dramatically while the consumption (eating) reward $\theta_{consumption}$ drops significantly. In both cases reward coefficients θ_4 reach the maximum value of 1, and this happens in $\sim 8,000$ s for weaker vitamins, versus a considerably faster $\sim 4,000$ s for stronger vitamins. These relative rates are also seen in the reduction of $\theta_{consumption}$ values. Conversely, θ_4 reduces to near zero in the ‘coins as poison’ condition, and coin collection also falls to very low levels (Appendix A.3). This is accompanied by an increase in $\theta_{consumption}$. In all cases, Ent populations remain viable throughout the experiments. The key observation from these results is that both reward coefficients and coin collection rates *increase* when coins are (at least superficially) beneficial (vitamins), and both *decrease* when coins are harmful (poisons).

Experiment 4: Dormant reward Figure 8 shows results from Experiment 4 (in which coins are absent from the environment during baseline training, and are then introduced at a

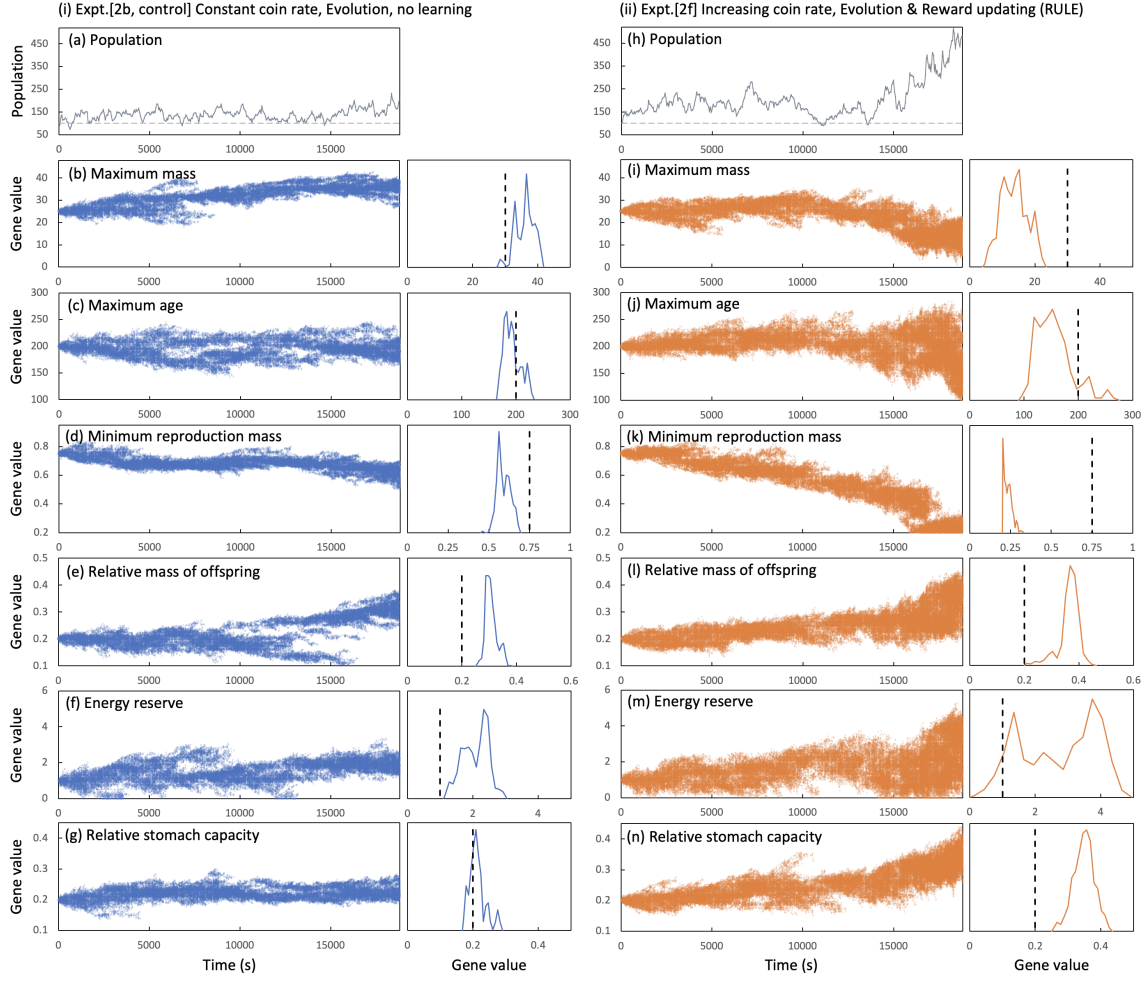


Figure 6: Evolution of gene values. Each point in the time series represents the gene value of an Ent, plotted at its time of birth. The gene value distributions are for the last 500 Ent births of the experiment, with the vertical dashed line indicating the initial gene value in each case.

linearly-increasing rate during the experiment; RULE is enabled and there is no evolution). Figure 8 parts (iii) and (iv) together show that θ_4 and the rate of *coin* collection remain consistently around zero for the 20,000 s of the experiment; conversely, θ_4 values and the rate of *vitamin* collection per Ent show significant increases over time after $\sim 2,000$ s. The upper limit of $\theta_4 = 1$ is reached at $\sim 15,000$ s and is associated with a stabilised collection rate. The difference in behavioural response to introduced coins and vitamins is indicated in Figure 8: by the end of the experiment, Ents are positioning themselves roughly evenly within the area where vitamins are dropped and collecting them quickly (ii), while avoiding the area where coins are dropped (i), and showing consistent disinterest in collecting them.

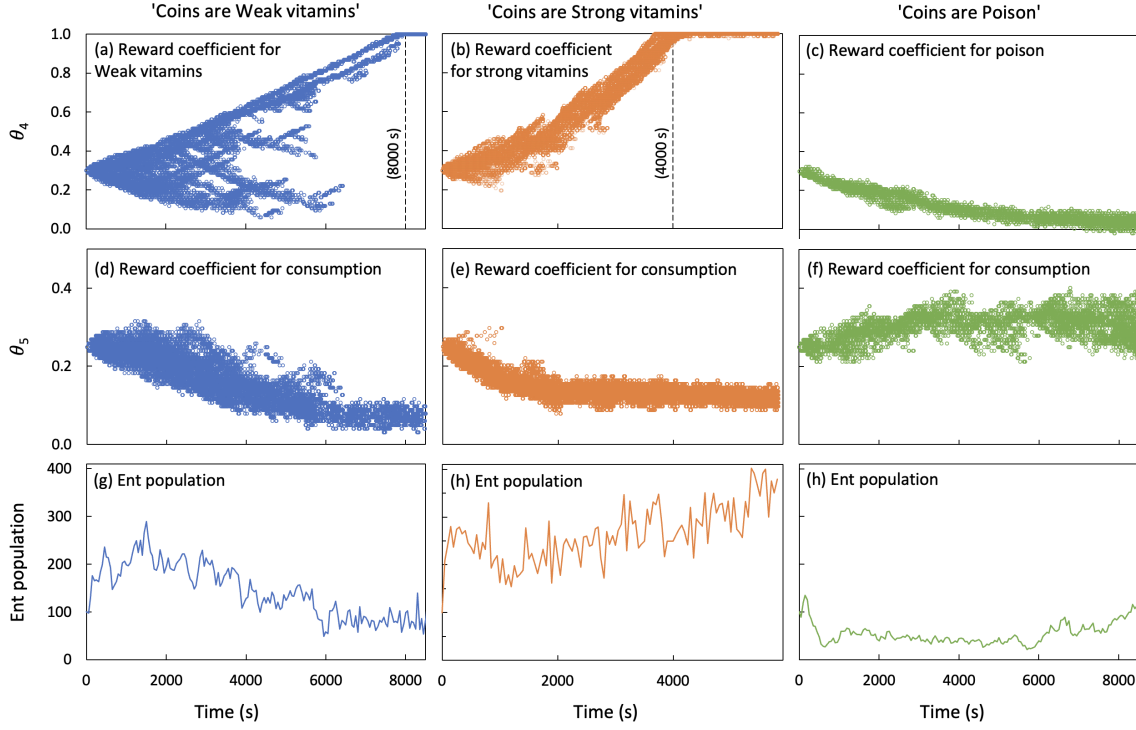


Figure 7: Vitamins and Poison. Results from three experiments in which ‘coins’ were recast as ‘vitamins’ or ‘poison’. Picking-up vitamins provides an instantaneous increase in stored energy and an increased probability of becoming reproductive. Conversely, picking up ‘poison’ reduces both stored energy and reproduction probability. The results shown are for cases where RULE updating of reward coefficients is enabled. The positive/negative benefits/costs of vitamins/poisons are reflected in the increased/decreased reward coefficients for ‘coins’ (θ_4).

Currencies 1-3 and Signals transfers Currency and signal transfers are not the core focus of the current experiments, and only brief mention is given here. Transfers of currencies between Ents are observed in the results, and signals were also passed. However, there were no major consistent trends in the data across the various experiments, suggesting these behaviours were not instrumental in surviving the coins challenge, or in surviving the otherwise benign conditions of the baseline experiments, and for the sake of brevity are not reported further.

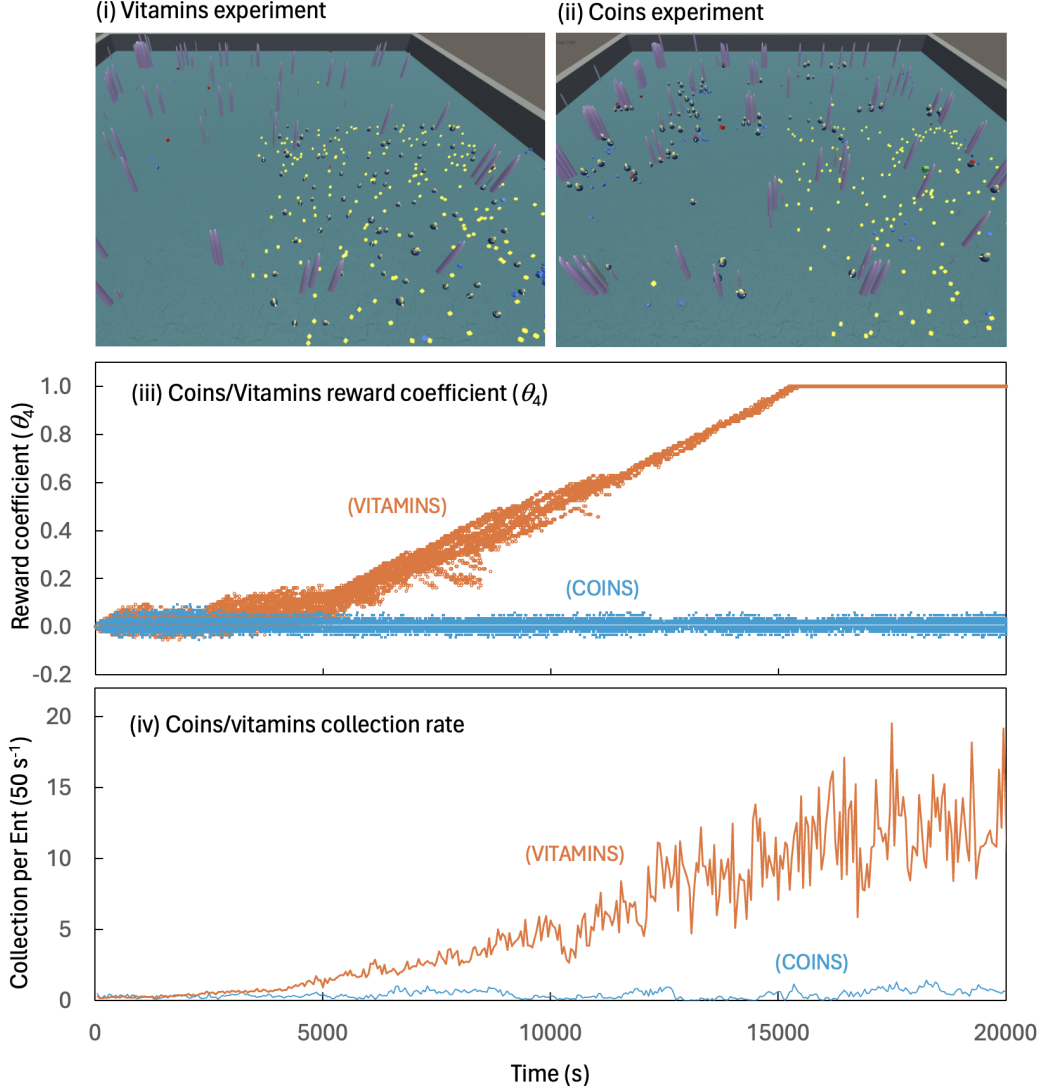


Figure 8: Dormant reward response to Coins and Vitamins. (i) and (ii) show example snapshots of the distribution of Ents in the environment during each experiment. The clustering of Ents close to Vitamins and the avoidance of Coins is observed. In (iii), each point represents the θ_4 value of an Ent born at the time indicated. (iv) shows the rate (per Ent) at which coins/vitamins were collected over the duration of the experiment.

6. Discussion

Death by coins Collecting coins is intentionally frivolous as it wastes time and energy that could be spent on more productive things (eating and reproduction, for example). Baseline coin rewards were set relatively high at the outset, to promote coin collection, with a small and fixed supply rate of coins promoting competition amongst the Ents (as was observed). With fixed rewards, high rates of coin supply are fatal to the Ent population: greater coin

availability per Ent means more time collecting them, hence the sometimes sharp populations collapse as more coins are collected by ever-decreasing numbers of Ents. Behavioural policies that would have resulted in higher rewards (with the same reward function), e.g. collecting *fewer* coins in order to preserve life (hence avoiding the -1 reward for early death) plus eating and reproducing more, were not discovered during training before populations collapsed. With large numbers of repeated attempts, such policies may have been discovered, but doing this in linear time adds additional (possibly insurpassable) difficulties. Indeed, if this were possible a single high-level sparse reward may indeed meet the challenges posed in this work (although this of course does not avoid the issue of choosing what that high level reward should be). Results show no evidence that (continuous) learning, with fixed reward coefficients, can solve the coins problem, as learning is always in service to the same reward function.

Testing RULE: the coins challenge The RULE algorithm is proposed as an approach to address the problem of fixed reward coefficients: parents (mothers) modify the values of reward coefficients and expected reward distributions passed on to their offspring, based on their own experiences. During continuous learning across the population, the policy is updated to maximize rewards and the modifications parents make to θ act to encourage their offspring to behave more like the parents. A diversity of experiences across the Ent population, and hence differences in the ‘success’ of individuals, is exploited to promote behaviour which is good for the population. The process is a dynamic majority vote to reinforce beneficial modifications to reward coefficient values, but also to leave open the possibility of reinforcing different behaviours. For example, in results for the second half of Experiment 2, the average θ_4 value is close to zero, but small sub-populations continue to have different coin collection experiences and produce short-lived ‘exploratory’ lineages, as excursions from the main trend. These excursions die out if the associated experience is not beneficial but it is this exploration of rewards that is needed to find new behavioural solutions to novel conditions (as seen in Experiments 3 & 4.). It is possible that perturbing the reward function in the way described may address potential issues related to loss of plasticity under attempted continuous learning tasks, as Ents appear to have little problem adapting their behaviour to novel environmental conditions when RULE is applied. The evidence supports the conclusion that dynamic beneficial reward updating is possible. This is true for coins, even though in themselves they are not harmful, and their impact on Ents is indirect via opportunity costs. RULE should work equally well, or even more ‘purposefully’ when faced with more direct impacts. To this end, Experiment 3 (‘vitamins and poisons’) was focused on understanding how the nature of the response to collectable objects depends on their properties.

Testing RULE: vitamins and poisons Vitamins provide clear benefits to Ents (increased energy and reproduction rate), their collection makes sense, and the results show as vitamins are made available, RULE increases θ_4 ; further, the associated increase in collection rate which follows reflects the strength of their benefits. The response to stronger vitamins is faster and more coherent, and there is little ‘exploration’ of lower θ_4 values (Figure 7(b)). For weaker vitamins, where the benefits are more marginal, there is more ‘exploration’ of lower θ_4 values before higher rewards eventually win out (Figure 7(a)). A greater level of ‘deliberation’ in less straightforward cases is perhaps inevitable given the presence of

conflicting reward-updating signals. In both cases θ_4 values reach their maximum limit of 1 and there is a major change in Ent behaviour. The Ent's consumption of PPs and processed material drops significantly, as they become reliant on vitamins (reward coefficient $\theta_{consumption}$ values drop significantly in both cases). This behavioural response to vitamins represents a major departure from the baseline behaviour of the Ent's.

The opposite response is seen in Figure 7(c,f): reduced θ_4 and increased $\theta_{consumption}$. Poisons have a negative impact on Ent energy levels and reduce reproduction rates. Under RULE, they are almost entirely abandoned (very few poisons are picked up) and Ent's navigate between/away-from poisons if they stray close them. The 'Poison' challenge is a difficult challenge as Ent's begin the experiment with strong poison(coin)-collecting behaviour (relatively high reward coefficients and associated policy) and therefore impose increasing levels of harm on the population as the poison (coin) supply increases over time. There is essentially a race between the harm inflicted (decreasing the population) and the adaptation of behaviour (increasing the population). The faster of the two poison (coin) increase rates exceeds the ability of the population to adapt (none survive), while the slower increase allows enough time for behaviour to successfully adapt and the large majority of populations survive (Table 3). The difficulty of this experiment, even for slower rates of coin increase, is seen in Figure 7(h) as a relatively low population. Low populations are at the mercy of stochastic variation and this is reflected in the lower proportion of populations surviving this experiment (75% versus 80% in Experiment [2e]). In summary, and as seen in results for the 'coins challenge', the diversity of experiences across the Ent population is exploited by RULE as it responds to both beneficial (collecting vitamins) and harmful (collecting poisons) behaviours, by appropriately modifying rewards and therefore learnt behavioural responses.

Testing RULE: dormant rewards The reward function components used in the present simulation, while defensible in that they produce persistent populations, are somewhat arbitrary, and this inevitably limits generalisability: only the reward components present can be updated and these may not necessarily apply to future changes in the environment. Two options to tackle this problem are discussed: (1) Predefined 'dormant' rewards, (2) Dynamic reward function expansion.

Option (1) involves increasing the fixed number of components in the reward function, with a subset of these being redundant at first (in the sense they contribute nothing to the total reward as the phenomenon they reward is absent and the coefficient is set to zero). In this case, at some point in the future RULE might be provoked by the environment in to 'switching-on' the rewards (increasing the absolute value of the coefficient), thus having it contribute more or less to the ongoing learnt behaviour. Option (2) calls for a mechanism to recognise new 'rewardable' phenomena and to dynamically add relevant components to the reward function (for example, when phenomena are above some threshold level of surprise). Option (2) approaches Option (1) as knowledge of possible future rewards increases. The question of how to practically implement option (2) is not addressed in this work.

Experiment 4 provides a test of how well RULE responds in the presence of a dormant reward (Option 1), in this case for coins and vitamins. When a persistent Ent population experienced coins for the first time (with $\theta_4 \approx 0$), a small number of coins were picked-up by chance. As coin availability was increased (linearly) over time, coins continued to be picked-up only at a low rate and rewards did not change significantly: coin collection was

not ‘switched-on’ to any significant degree. Coins provided no benefit. These findings are consistent with those of Experiment 2. Conversely, after vitamins were first experienced, the dormant reward showed signs of being ‘switched-on’ after ~ 1 ks, with rewards and collection rates increasing significantly (θ_4 eventually reaching its maximum value). These observations suggest RULE has the potential to respond appropriately to novel conditions, setting reward coefficients to values which reflect the benefit of the action. These results underscore how ‘artificially’ high the baseline value for θ_4 was set, and that given the chance to either start at $\theta_4 = 0$ and discover coins for themselves, or to start at $\theta_4 = 0.3$ and adjust to the reward, Ents using RULE settle on $\theta_4 \approx 0$.

Impacts of Evolution With broadly stable conditions, no evolution of the primary producers, and only a single species of Ent, there is limited evolutionary pressure in the present set of experiments. Nonetheless, if the initially chosen baseline gene values were sub-optimal (which is likely), evolution would be expected to fine-tune these values over time, and this is essentially what is observed. Each of the consistent gene-value changes observed under baseline conditions can be interpreted as aiding survival (larger Ents with greater energy reserves who start to produce larger offspring earlier in their lives). Under the coins challenge experiment, evolutionary changes alone were not sufficient for the population to survive, but in combination with learning and reward updating (under the slower of the two coin increase rates), systematic changes in gene values did produce Ents with characteristics well-suited to surviving high-coin-rate conditions, and resulted in the largest populations recorded across all repeated experiments. However, as shown in Figure 5(1), imposing the higher rate of increase in coin availability caused the population to collapse (results which were consistent across multiple repeat runs). In ecological settings, populations may experience a lag in adaptation when conditions change rapidly, meaning relatively slowly-changing traits which were once beneficial become harmful under new conditions. In the present case, population collapse happens before large systematic changes in mean gene values have happened, but there is a significant increase in the *spread* of gene values ($\sim 25\%$ mean increase in relative standard deviation). There is a potential asymmetry present in that random combinations of gene values which aid survival have relatively small effects (as seen in Experiment 2a,b), whereas those which hinder survival have enough of an effect, in the presence of high coin-stress, to reduce the population. Once the population starts to decline, positive feedback (increasingly higher availability of coins per Ent) takes over (as discussed above). It appears then that if RULE can provide resilience to coins, Evolution fine-tunes the Ents, but if the increase in environmental stress is too fast, the negative aspects of gene value variation collapse the population before RULE can save it.

Reward hacking The goal of RL is to tune a policy to maximizes long-term rewards, and during continuous learning the RULE algorithm modifies reward coefficients based on policy-driven behaviour. In principle then, the training process could hijack the adjustment of actions in order to drive a modification of rewards, thus boosting the rewards obtained (reward hacking). In fact, multiple simultaneous interacting optimisation processes are at work in some experiments: implicit evolutionary optimisation of gene values, explicit maximizing of rewards through policy updating, and the RULE-based reward coefficient updating. Potential exists for any of these to amplify or dampen each others’ impacts. Constant baseline conditions, with RULE and evolution enabled, and all but one (reproduction) of the reward

coefficients set below their maximum value of 1.0, provides a good base from which to test for reward hacking. In the results from this case (the control condition of Experiment 2f), no systematic increase in reward coefficients is observed. It is possible that survival needs impose strong constraints on what behaviours can be rewarded (increasing rewards for one activity potentially means less of another), and that this keeps reward hacking in check. There are however other observations which paint a more nuanced picture. During the coin challenge (Expt. 2e; RULE with no evolution), θ_4 is reduced to a level where coin collection still happens, but at a much-reduced level. That is, the reward from coins is not lost, but is reduced to a point which does not threaten the population (in stark contrast to the case of weaker poisons, where collection was halted altogether). The response to Experiment 2e created a population largely un-interested in coins, living outside the area where coins were dropped, with a small fraction of the Ents collecting coins at relatively low rates in the coins area. When evolution is added to this condition (Expt. 2f), the body characteristics of Ents are changed significantly, resulting in a very large population of shorter-lived Ents who as a population are able (by the end of the experiment period) to collect very high levels of coins without the population collapsing. This ‘solution’ to the coins challenge is therefore successful in both producing a surviving population and also maximising the reward from coins. This is perhaps more of a ‘reward co-operation’ between processes than a ‘reward hacking’ by one of them. It is possible that if the complexity of the environment was increased further, agents may find other opportunities to hack rewards, but the constraints imposed by survival (in the linear timeframe) may dampen such tendencies. This remains an open question and an avenue for further work.

Hidden goals In the introductory sections it is argued that the goals of autonomous real-world agents are typically hidden, and that survival challenges guide the emergence of rewards to satisfy those hidden goals. Since RULE can adjust reward coefficients, it is natural to consider whether the resultant reward function might be interpretable in terms of the hidden goal(s) it serves. It is argued below that this is not possible. Updated rewards elicit individual behaviours which result in emergent consequences at the population level (as in the present case, agents operate within the context of other agents). To take an extreme hypothetical case, imagine Ents using RULE see an increase in the reward for consumption, which rises to the maximum value of 1. At the same time, all other rewards fall to low levels. Interpretation of the goal may appear simple: maximizing consumption of food is the most important factor in survival. However, when all of the population focusses on eating, there is a secondary effect such as out-competing a second species which would otherwise have competed in some other way (e.g. competition for space). The imperative driving the true goal therefore is reducing competition for space. This was achieved in the present case through competing for food, but could feasibly have been achieved in other ways. This is a relatively simple case and yet the true high level goal is not easily derived from the reward function, or at least not uniquely. When there are multiple reward components and more complex behaviours, it seems likely identifying goals would become more difficult. Hidden goals therefore likely remain hidden, even when the results of reward function updating are known.

Wider application of RULE To what extent can RULE be used in other multi-agent contexts? RULE operates at the population level by exploiting variation in individual experi-

ences, and works essentially because successful/unsuccessful behaviour is amplified/dampened. The current context approximates a simple ecosystem, where entities face a constant survival challenge. One potential solution to generalising RULE is to cast other situations as analogues of the survival challenges of individuals. For example, economic agents may have money in place of energy, costs such as wages instead of base metabolism, acquisitions as foraged food, sales as consumption, expansion as reproduction, bankruptcy as death, etc.. On social media platforms, influencers and content creators vie for limited user attention and engagement, where audience size mirrors reproduction. While such analogies can be somewhat strained, RULE is expected to work well in situations where populations of decision-making entities exist within resource-constrained environments (defined broadly), where there is scope for differing experiences which affect outcomes driving implicit or explicit competition. The action space of entities in the current environment is intentionally relatively small and it remains to be seen whether the RULE algorithm can indeed scale to even more complex environments, e.g. Baker et al. (2022). A useful further benchmarking exercise would be a comprehensive comparison of RULE to RL approaches based on fixed higher-level intrinsic rewards (such as novelty search) and possibly hybrid schemes which combine aspects of these approaches with RULE. The RULE approach should in principle also apply in cases where entities are not individuals, but collectives of individuals. For example, in supply chain networks, manufacturing firms, distributors, and retail companies, are decision-making entities which compete for resources and customers under constraints of costs. Similarly, city entities compete for population, funding, and natural resources, with expansion through infrastructure development and decline through population and economic loss. Where individual agents in the simulated system are not of a type which produce offspring (or have enforced maximum lifespans), a different mechanism is required. Agents could, for example, periodically replace themselves with updated versions (effectively reproducing ‘in place’) when some resource condition was met, with calculated adjustments to θ_i and $E(\tau)$ based on comparisons of accumulated rewards to expected rewards over a rolling time window.

7. Conclusion

Central to this work is the idea that goals and reward functions of real-world entities, which we may wish to represent in models, are hidden from observers (and most likely also from the entities themselves). Further, the reward functions that guide behaviour are in some sense chosen, or emergent, in real-world systems and adapt over time as environments changes. If out of a necessity to elicit some desired aspects of the entity’s observed behaviour, we formulate a fixed reward function, we risk pre-defining incorrect behaviour in novel situations for which we have no observational data, potentially limiting future adaptability. Hence there is a need to model entities which endogenously shape their own reward functions over time, particularly in response to changes in their environments, and thereby update their own behaviour for their own benefit. If our aim is to reproduce the dynamics of real-world systems, entities must also re-shape their behaviour in *linear time*, without the chance to learn by resetting their environments when they fail in some sense. The evidence presented from simulations of entities (Ents) within a simplified ecosystem-like environment shows that continuous Reinforcement Learning, together with continuous updating of reward functions using the RULE algorithm (Reward Updating through Learning and Expectation),

can achieve these goals. The algorithm exploits the diversity of experience across the Ent population to endogenously reinforce beneficial behaviours and reduce those which are harmful. This was demonstrated first by Ents ‘choosing’ to abandon previously learnt and positively-rewarded behaviours when they became harmful. Populations also responded appropriately to previously unseen objects, by creating positive rewards for (and collecting) beneficial ‘vitamins’, and holding the reward at zero (and not collecting) ‘coins’ with no intrinsic benefits when these were introduced in to their environment. The relatively simple ecosystem-like context of the present experiments was chosen as a ‘toy’ example because it contains generic properties found in many systems, including multiple competing agents, scarce resources, a spatial domain, costs, and exchanges of information. Applying RULE to yet more complex environments, in cases where multi-agent problems are cast as survival challenges, is potentially a useful focus for future work.

Appendix A. Additional model description

Broad descriptions of the simulated environment, its model components and their behaviours are provided in the main text. Finer details of aspects not covered there are provided below.

A.1 Primary Producers

Each Primary Producer (PP) updates its state every $P = 25\Delta t = 0.5$ s (the model update frequency is $50s^{-1}$; $\Delta t = 0.02$ s). The characteristics of each PP are determined by its gene values. In the present simulations, evolution is not enabled and when reproducing each PP simply passes on an exact copy of its genes.

Genes Each PP has its own set of genes which determine its characteristics, as outlined below.

Description	Symbol	Units	Range	Initial value
G[0] maximum biomass	m_{ppMax}	kg	> 0	50
G[1] ‘Lock’ L1 [colour, r]	L_1	-	$[0, 1]$	0.9
G[2] ‘Lock’ L2 [colour, g]	L_2	-	$[0, 1]$	0.7
G[3] ‘Lock’ L3 [colour, b]	L_3	-	$[0, 1]$	0.9
G[4] Probability increment per 1% of accumulated mass	r_{inc}	-	$[0, 1]$	$5 \cdot 10^{-5}$
G[5] Initial reproduction probability	r_0	-	$[0, 1]$	0
G[6] Maximum age	a_{max}	s	> 0	100
G[7] Gene copying noise	z_{pp}	-	$[0, 1]$	0
G[8] Energy harvesting coefficient	$c_{s,m}$	-	> 0	2
G[9] Body mass energy density	ρ_{Epp}	E/kg	> 0	5
G[10] Maximum reproduction probability (per dt)	r_{ppmax}	-	$[0, 1]$	0.005
G[11] Exclusion radius to height ratio	h_{ex}	-	> 0	3
G[12] Growth rate (kg growth per kg of body mass per s, when s=1)	g_{pp}	s^{-1}	> 0	0.5
G[13] Pre-growth pause	t_{pause}	s	> 0	5
G[14] Inner exclusion ratio	r_{in}	s	> 0	0.2
G[15] Proportion of seeds on surface	p_{surf}	-	$[0, 1]$	0.05

Table A.1: Gene values for Primary Producers. Gene index numbers are shown. Genes [1,2,3] are jointly the {r,g,b} colour tuple (expressed as ratios) for the PP outward appearance and the ‘lock’ values L1-3 which determine palatability. As PP evolution is not enabled in the present case, the initial values shown remain fixed throughout all model runs.

Reproduction via seed production Each PP produces a fruit within any given update step with reproduction probability r , which is initialised at r_{pp0} and increases over time if $r_{pp} < r_{ppmax}$ where r_{ppmax} is a genetically-set maximum reproduction probability. The increment increases to a maximum, as

$$\Delta r_p = \frac{m_{pp}}{m_{ppMax}} \cdot r_{inc} \cdot p + n_{bites} \cdot r_{inc} \quad (A.1)$$

where, m_{pp} is current PP mass (kg), $m_{ppMax} = 50kg$ is the genetically-set maximum mass, $r_{inc} = 5 \cdot 10^{-5} dt^{-1}$ is the reproduction probability increment, n_{bites} counts the number of times the PP has been bitten by Ents since the last fruit production. This provides feedback

from Ent behaviour to PP behaviour, relevant to co-evolution of Ents and PPs, but has limited impact at present as evolution of PP gene values is not enabled in the simulations reported.

When fruiting occurs, fruits are dropped at a random position between an inner and outer calculated radius, with radii satisfying $\max(h \cdot r_{in} \cdot 0.5, 0.3)$, where, $h = 0.005m_{pp}$ is the height of the PP and $r_{in} = 0.2$ is the genetically set inner exclusion ratio.

A genetically determined proportion ($p_{surf} = 0.05$) of the seeds produced remain on the surface for a period of 10 s, and then transition below the surface. During their period at the surface seeds can be picked up and stored by Ents (details below). After a genetically determined ‘germination’ period of duration $t_{pause} = 5$ (s), seeds begin to grow symmetrically out from the surface and deeper in to the ground.

Growth Condition When illuminated, PPs add biomass in increments Δm_{pp} while $m_{pp} < m_{ppMax}$.

$$\Delta m_{pp} = E_s \cdot c_{s,m} \cdot m_c \cdot \frac{g_{pp}}{\rho_{E_{pp}}} \cdot \Delta t \cdot P \quad (\text{A.2})$$

where $E_s = 1.0$ is the energy input from the external source, $c_{s,m} = 2$ is the external energy collected (collection area \times conversion efficiency), $g_{pp} = 0.5$ is a growth constant (kg growth per s, per kg of biomass), $\rho_{E_{pp}} = 5$ is the body mass energy density.

Competition for space To ensure some degree of spacing between PPs, PPs mimic the behaviour of many real-world primary producers and exert direct competitive pressure on their neighbours. A PP competes with another PP if it is within its inner (r_{inner}) and outer (r_{outer}) competition radii. If they compete, the larger of the two competing PPs has a 10% chance (per Δt) of killing the smaller PP.

$$r_{inner} = h \cdot \zeta_x \quad (\text{A.3})$$

$$r_{outer} = h_{max} \cdot \zeta_x \quad (\text{A.4})$$

where $h_{max} = 0.005 \cdot m_{max_{pp}} = 0.25m$ is the maximum PP height and $\zeta_x = 3$ is the genetically-set exclusion-distance-to-height ratio.

Death PPs die if (i) they reach the genetically-determined maximum age ($G[0]=100$ s), (ii) they are out-competed (as outlined above), or (iii) they are consumed to the extent their biomass is reduced to zero. Upon dying, the PP does not instantaneously disappear, but reduces its remaining mass linearly at a rate $-m_{fade}$.

A.2 Ents

A.2.1 ENERGY

Energy available to Ents is obtained from consumption and can be stored within their bodies, converted to biomass (growth), used to meet basic metabolic costs and to undertake actions, all of which have an associated energy cost. Energy is quantified in arbitrary ‘energy units’ and referred to inter-changeably as currency-zero (C_0).

Metabolic Cost Base metabolic cost is subtracted automatically.

$$k_{base} = \nu \cdot b \cdot L^{0.667} \quad (\text{A.5})$$

where, k_{base} is base metabolic cost rate, L is the lean mass of the entity, I is the ratio of infrequent updates, used to adjust the calculation based on the frequency of updates.

Consumption If an Ent is touching an object, it automatically evaluates whether the object is a soft and therefore whether it can be bitten (Ents, PPs and processed blue-cakes are soft). If the Ent chooses to bite the object, it consumes some of its mass, which is immediately removed from the other object. The Ent obtains energy by digesting the consumed mass, the amount of which depends on multiple factors, including the mass, energy density and palatability of the consumed material.

$$m_{eaten} = \min(q_{max} - q, m_{obj}) \quad (A.6)$$

where, m_{eaten} is the mass consumed, q_{max} is the maximum stomach capacity (calculated as $m_{ent} \times q_f$, where m_{ent} is the lean bodymass of the Ent (see below) and q_f is the fractional stomach capacity), q is the current mass in the stomach, and m_{obj} is the mass of the object. The mass consumed is digested over time in a series of increments, and as each increment is digested the Ent receives energy

$$\Delta q = (q_{max}/50t_{dig}) \quad (A.7)$$

where, Δq is the digestion mass increment, q is the stomach content, and t_{dig} is the digestion time needed.

$$t_{dig} = q/(q_{max} \cdot d_r) \quad (A.8)$$

where d_r is the basic rate at which the stomach can process its contents. The energy derived from each digested increment is calculated as

$$\Delta C_{0_{con}} = \rho_{Ent} \cdot \xi_{con} \cdot \Delta q \cdot \eta \quad (A.9)$$

where, $\Delta C_{0_{con}}$ is the energy available, ρ_{Ent} is the energy density of the consumed material (E/kg), ξ_{con} is the conversion efficiency for consumed mass, and η is the nutritional value coefficient.

For processed blue-cake, the nutritional value is set as $\eta = 1$. For other consumed material, η depends on the *palatability* of the consumed material, which is determined by ‘key’ $\{K_1, K_2, K_3\}$ values of the consumer Ent and the ‘lock’ $\{L_1, L_2, L_3\}$ values and of the Ent or PP being consumed.

Palatability (P) is based on genetic compatibility over the lock and key values

$$P = \frac{\sum_i 1 - (K_i + L_i)}{n} \quad (A.10)$$

with $n = 3$ in this case. The nutrition coefficient is bounded within $[-1, 1]$, calculated as

$$n_v = \max(\min(1 - 4 \cdot |p|, 1), -1) \quad (A.11)$$

where $|\cdot|$ is the absolute value.

Energy partitioning After energy is obtained from digestion, it is routed to either biomass growth or storage (for later use) via a growth-to-free-energy ratio g . Ents maintain a

reserve of energy, as a buffer against scarcity, and the size of this reserve (E_{res}) is controlled by gene value $G[34]$. The partitioning ratio g is determined by:

$$g = \begin{cases} 0 & \text{if } L \geq m_{entMax}, \\ \max(0, \min(1, g_0 \cdot (1 - m_{ent} \cdot g_{con}))) & \text{if } L < m_{entMax} \text{ and } C_0 < E_{res}, \\ 1 & \text{if } L < m_{entMax} \text{ and } C_0 \geq E_{res}. \end{cases} \quad (\text{A.12})$$

Here, m_{ent} is the lean mass of the Ent, m_{entMax} is the maximum lean mass ($G[0]$), C_0 is the amount of energy being stored (as fat) by the Ent, E_{res} is the energy reserve level, $g_0 = 0.5$ is the initial value of g ($G[8]$), g_{con} is the growth-energy constant defined by

$$g_{con} = g_{\Delta max} / m_{entMax} \cdot g_0 \quad (\text{A.13})$$

where, $g_{\Delta max} = 0$ in the present case, meaning the second condition in Eq. A.12 reduces to g_0 .

This logic gives priority to filling the energy reserve; once this is full, energy is shared equally between growth and additional storage until growth is complete; then, energy obtained from digestion is all sent to storage for later use.

Growth Growth occurs incrementally once each 0.25 s.

$$\Delta m_{ent} = \Delta C_{0con} \cdot g / \rho_{Ent} \quad (\text{A.14})$$

where ρ_{Ent} is the energy density (E/kg) of the Ent.

Starvation If an Entity does not eat, it continues to operate by using its energy reserve. If its reserve gets depleted, biomass is converted directly to energy (equivalent to catabolism in biological life) at a rate

$$\Delta C_{0con} = \Delta m_{ent} \cdot \psi \quad (\text{A.15})$$

where $\psi = 0.9$ ($G[47]$) is the conversion efficiency between Ent biomass and energy. This process can continue until the Ent's body mass (m_{ent}) reaches a minimum viable level, after which the Ent dies. This 'point of starvation' is dynamically updated, being a fixed fraction of the largest previously-achieved lean biomass.

$$m_{starv,t} = \max(m_{starv,t-1}, m_{ent} \cdot f_{starv}) \quad (\text{A.16})$$

where, $m_{starv,t}$ is the current minimum viable body mass, $m_{starv,t-1}$ is the value until the previous time step, m_{ent} is the lean mass, and f_{starv} is the minimum viable body mass fraction.

A.2.2 MOVEMENT

Ents can choose to move, which involves deciding on rotation and applied force. The simulation is physically based and so other factors also influence Ent movement such as collision with other (potentially moving) objects, friction (both linear and rotational) acting against the applied forces, and gravity.

In each time step (0.02 s), Ents update their rotation by choosing an action $a_{rot} = \{-1, 0, 1\}$, which defines their change in rotation $\Delta\Phi$.

$$\Delta\Phi = \Delta t \cdot t_{rot}^{-1} \cdot a_{rot} \cdot 360 \quad (\text{A.17})$$

where, t_{rot} is the time required for a single rotation and is set by gene G[23], a_{rot} is the chosen rotation action (either of $\{-1, 0, 1\}$), and the value 360 converts the rotation to degrees.

Similarly, Ents choose how to increment their velocity by applying force F in either a forwards or backwards direction aligned with their rotation angle.

$$F = \max(F_{max} \cdot m_{ent} \cdot (1 - q_f), 100) \quad (\text{A.18})$$

where F_{max} is the maximum force per kilogram the Ent can apply (set by gene value G[9]), m_{ent} is the lean mass, q_f is the stomach mass ratio. The maximum velocity of Ents ($v_{max} = 1$ m/s) is set by gene value G[46]. A minimum value of $F = 100$ is enforced so that even small Ents can overcome friction and move. The inclusion of $(1 - q_f)$ means there is a trade-off when evolution is enabled between larger stomach size (ability to consume faster) and ability to move quickly. The total mass of the Ent (which the applied force is acting on) includes lean mass, fat mass, stomach content and also any items being carried, and the force in proportion to net lean mass represents the force generating part of the Ent body (muscle, ligaments, bone).

Movement cost Generating force for movement costs energy and this is calculated as k_{move} based on the magnitude of force applied and a predefined rate of energy consumption.

$$k_{move} = v_k \cdot F \quad (\text{A.19})$$

where, $v_k = -4 \times 10^{-5}$ is the energy consumption rate per unit of force. An amusing mistake during the early phase of this work led to the sign of velocity being reversed in the movement cost calculation. During training, Ents quickly learnt they could maximize their energy inputs, without going to the trouble of eating, by travelling backwards as fast as possible in tight circles.

Collision damage Movement can also be associated with a different cost, that associated with collision damage. For some sense of realism, Ents should learn not to collide at high speeds with solid objects. The impact damage calculation models the collision of two objects based on their kinetic properties. Assuming the position vector of the first object is \vec{p}_1 and the point of collision is \vec{c} , the vector \vec{d} , pointing from the position of the first object to the point of collision, is calculated as

$$\vec{d} = \vec{c} - \vec{p}_1 \quad (\text{A.20})$$

The velocity component along the collision direction is given by

$$v_{line} = \vec{d} \cdot \vec{v} / \|\vec{d}\| \quad (\text{A.21})$$

where \vec{v} is the velocity vector of the object, and $\|\vec{d}\|$ is the magnitude of the collision direction vector. The effective mass used in the damage calculation is

$$m_{eff} = \min(10m_1, m_2) \quad (\text{A.22})$$

where m_1 is the mass of the first object and m_2 is the mass of the second object. This formulation limits the impact of extremely disparate mass values on the calculation. The relative impact damage is calculated by

$$D_{rel} = -(\max(0, v_{line}) \cdot m_{eff} \cdot k) / (m_1 + m_{eff}) \quad (\text{A.23})$$

Here, k is a coefficient representing the impact damage sensitivity, which scales the damage based on the physical properties and the dynamics of the collision. The use of $\max(0, v_{\text{line}})$ ensures that only impacts where objects move towards each other are considered, avoiding negative damage values from receding collisions. The total impact damage inflicted by the collision is then computed as

$$D = D_{\text{rel}} \cdot m_1 \cdot \psi \quad (\text{A.24})$$

where ψ is the same conversion efficiency between biomass and energy as described above in the context of starvation.

A.2.3 REPRODUCTION AND INHERITANCE

Ents do not have different sexes, and each can act as both a ‘mother’ (which gives birth to offspring) and a ‘father’ (which is required to pass on their genes to mothers, triggering reproduction). Ents can enter the ‘mother’ state if the following conditions are satisfied:

$$m_{\text{ent}} \geq m_{\text{repMin}} \quad \text{and} \quad C_0 \geq E_{\text{birthTotal}}, \quad (\text{A.25})$$

where m_{ent} represents the lean mass of the entity; m_{repMin} is the minimum mass required for reproduction; C_0 is stored energy; $E_{\text{birthTotal}}$ is the total energy cost required to produce an offspring. This total energy cost is met by the mother and is defined

$$E_{\text{birthTotal}} = -(n_{\text{offspring}} \cdot m_{\text{offspring}} \cdot \rho_{\text{ent}}) - C_{0,0} \quad (\text{A.26})$$

where $n_{\text{offspring}}$ is the number of offspring produced (fixed at 1 in the present case), $m_{\text{offspring}} = m_{\text{entMax}} \cdot o_{\text{relMass}}$ is the mass of each offspring ($o_{\text{relMass}} = G[19]$ being the genetically-determined relative offspring mass), and $C_{0,0}$ is the energy given by the mother which goes in to the offspring’s (fat) store (the mother also contributes initial amounts of currency C_1 and C_2 equal to $C_{\text{initial}} = 0.1$; G[43]). Upon satisfying the conditions of Eq.A.25 the probability of entering the mother state (per 0.25 s) is p_{rep} , where

$$p_{\text{rep}} = \nu_r \cdot (m_{\text{repMin}}/m_{\text{entMax}})^{-1} / (\nu_{\text{global}} \cdot a_{\text{max}}/\lambda_{r\text{Max}}) \quad (\text{A.27})$$

where $\nu_r = 25$ is the timesteps between each reproductioni state update, $\nu_{\text{global}} = 50$ is the global update frequency (s^{-1}), $\lambda_{r\text{Max}}$ is the genetically-determined expected number of offspring produced per maximum Ent lifetime.

Once a ‘mother’ Ent is in the reproductive state, in order to produce an offspring a ‘father’ Ent must physically touch the ‘mother’. When this happens, the genetic distance between the two entities (Δ_g) is used to assess their compatibility:

$$\Delta_g = \frac{\sum_{i=1}^3 |g_{1i} - g_{2i}|}{3} \quad (\text{A.28})$$

where, g_{1i} is the i th gene of the first entity, and g_{2i} is the i th gene of the second entity. Reproduction is successful if $\Delta_g < \Delta_{g\text{max}}$, with $\Delta_{g\text{max}} = 0.15$ being the maximum allowable genetic difference for reproductive compatibility. This value ($\Delta_{g\text{max}}$) is itself determined by a gene value, G[42]. If reproduction is allowed, the offspring’s gene values are a random

selection of its parents’ genes (with each gene value having a 50% chance of coming from either parent). Once selected, each value is perturbed by noise, according to

$$G[i] = G[i] + \delta_{G[i]} \cdot \tilde{U} \quad (\text{A.29})$$

where $\delta_{G[i]} = 0.015$ is a ‘copying noise’ parameter, determined by gene value $G[56]$, which means the rate of evolution is to some extent controlled by evolution; \tilde{U} is a uniform random number over the range $[-1,1]$. Upon successful reproduction, the Ent offspring appears in the simulated environment next to the mother Ent. It then contributes observations, actions and rewards to the learning process along with other Ents.

A.2.4 REWARD UPDATING

The RULE algorithm is outlined in the main paper. Constants α_i and β_i (where i is reward coefficient number) are calculated as follows: run simulation under background conditions with chosen θ_i ; while running, record rewards accumulated at each time bin τ for each Ent, and continue simulation until mean accumulated reward in each age bin are stable, yielding expected values $E_i(\tau)$; set $T_i = \sum_{\tau} E_i(\tau)$, $n_i = T_i/\theta_i$, $\alpha_i = T_i/n_u$, where $n_u = 25$ was user-defined, reflecting the number of generations over which reward adjustment occurs (n_u could of course be added to the gene to make this an evolved choice). Recall that α is the increment for updating $E_i(\tau)$ under RULE and β_i values increment θ_i . An unstable response (where θ_i values are always driven to either their minimum or maximum values) is observed when $\beta \geq T_i/n^2$, and so $\beta_i = c_i T_i/n^2$, with $c_i < 1$.

A.2.5 OBSERVATIONS

Observations are made automatically (Ents do not have to choose to make observations).

‘Vision’ Ents do not have access to images obtained from their point of view. This approach was tried in the early phases of development, and while it worked, it offered no advantages over the chosen approach, was more computationally costly, and produced slower learning. Therefore, observations analogous to vision are made by Ents using ray-casting within Unity. Notionally, 17 beams are sent from the Ent (a central beam at 0deg, and 8 equally spaced beams either side, providing an 80 deg field of view). Each beam is a cone, expanding away from the Ent to a sphere of radius 0.05 m. The length of each beam l_{ray} is determined by the strength of the external energy (light) source, so Ents ‘see’ further in stronger light.

$$l_{ray} = \max(1, -15 + 65 \cdot s_{light}) \quad (\text{A.30})$$

where s_{light} is the light intensity in normalised units, which for convenience is equal to the normalised energy source value E_s . Each beam returns data on whether the beam hit anything, what that object was (a one-hot encoding of an object tag list), and distance to the object. This information is updated each 0.02 s, and contributes to the total set of observation used to inform Ent behaviour and learning. In addition to this ‘vision’ data, 22 other observations are listed below in Table A.2.

A.2.6 ACTIONS

Picking-up objects Entities have the capability to interact with and pick up objects based on specific conditions. This mechanism is crucial for simulating realistic interactions and

Observation	Observation Description	Values
1	Normalized X velocity of rigidbody	[0, 1]
2	Normalized Z velocity of rigidbody	[0, 1]
3	Normalized Y rotation angle	[0, 1]
4	Starvation level	[0, 1]
5	Raw material presence	{0, 1}
6	Processed material presence	{0, 1}
7	Carrying cube	{0, 1}
8	Reproduction readiness	{0, 1}
9	Normalized currency 0	[0, 1]
10	Normalized C_1	[0, 1]
11	Normalized C_2	[0, 1]
12	Normalized C_3	[0, 1]
13	Normalized C_4	[0, 1]
14	Absence of broadcast signal	{0, 1}
15	Broadcast signal 1 detected	{0, 1}
16	Broadcast signal 2 detected	{0, 1}
17	Genetic distance of closest entity	≥ 0
18	Normalized distance to broadcast signal source	[0, 1]
19	Normalized angle to broadcast signal source	[0, 1]
20	Normalised number of entities within detection range	≥ 0
21	Detect being bitten/eaten	{-1, 0}
22	External energy source strength	≥ 0
[in addition, Vision information from 17 rays cast]		

Table A.2: Ent observations. In the table $\{\}$ indicate discrete sets of observations, $[]$ indicate continuous inclusive ranges. Each Ent collects and processes observations every 0.02 s. See text for information on vision observations. Some raw observed values are divided by a normalization factor: [Observations 1,2] Velocity is normalized by maximum velocity $G[46]$; [3,19] Rotation angle is normalized by 360; [4] The Starvation Level is defined $S = 1 - (m_{starv,t}/m_{ent})$; [9-13] All currency values are normalized by 100; [17] Genetic distance to the physically nearest Ent is calculated as Δ_g (Eq.A.28), and is referred to as a ‘sense of smell’ in the main body of the paper; [18] Distance is normalised to the sensing radius ($G[41]$); [20] The number of Ents within the sensing radius is divided by 100.

Action	Equation / Condition
1. Attempt to bite object	{0=False, 1=True}
2. Give currency	{0=False, 1=Give C_1 , 2=Give C_2 , 3=Give C_3 , 4=Give C_4 }
3. Synthesize currency	{0=False, 1=True}
4. Send signal	{0=False, 1=Emit_signal_1, 2=Emit_signal_2}
5. Update rotation	{-1=Rotate_anitclockwise, 0=No_change, 1=Rotate_clockwise}
6. Update speed	{-1=Apply_negative_force, 0=No_change, 1=Apply_positive_force}
7. Attempt pick up object	{0=False, 1=True}
8. Drop object	{0=False, 1=True}

Table A.3: Ent discrete action choices. Choices are made by each Ent every 0.02 s.

behaviours such as resource gathering or item collection. Even if an entity has the intent to collect objects from its environment, it can only pick up objects lighter than itself, simulating a basic physical constraint.

$$m_{object} < m_{entity} \quad (\text{A.31})$$

where m_{object} is the mass of the object being considered for pickup, and m_{entity} is the mass of the entity.

The entity’s storage capacity is equal to its (lean) mass, so new objects can only be stored when:

$$w_{storage} + m_{object} < m_{entity} \quad (\text{A.32})$$

where $w_{storage}$ is the current total weight in the entity’s current storage.

Training The proximal policy optimisation, PPO (Schulman et al., 2017) algorithm was employed for RL, implemented through the ML-Agents Toolkit within Unity (Juliani et al., 2020). A Python API provides a low-level Python interface for communication between the simulation and relevant PyTorch libraries. In the present case, only a single type of Ent is present in the simulations and PPO is implemented with a single policy, with all active agents contributing to the learning process and sharing that same policy. The following hyper-parameters were used during model training: hidden-layers=2x256, gamma=0.99 (discount factor for future rewards), batch-size=32 (number of experiences in each iteration of gradient descent), buffer-size=320000 (the number of experiences collected before updating the policy), time-horizon=32 (the number of steps of experience to collect per-agent before adding to the experience buffer), learning-rate=0.000025 (gradient descent rate, held constant), beta=0.0035 (strength of the entropy regularization, which ensures exploration by adding randomness to the policy; held constant throughout learning), epsilon=0.15 (a control on the evolution rate of the policy during training, controlling the divergence between current and updated policies; held constant constant during training), lambda=0.99 (a regularisation parameter, weighting the contributions from current and updated policy). During the initial training phase, the maximum number of steps was set to $1e8$ (recall there are 50 steps per second, and typically approx.100 agents contributing to the process, so this equates to a time limit of approx. 20,000 s). In subsequent experiments the maximum number of steps was set to infinity, and an upper time limit of 20,000 s was enforced.

Gene	Description	Symbol	Units	Range	Initial Value
G[0]*	Maximum biomass	m_{EntMax}	kg	[0.01, 50]	25
G[1]	Lock L1 & rgb[0]	L_1	-	[0, 1]	0.05
G[2]	Lock L2 & rgb[1]	L_2	-	[0, 1]	0.20
G[3]	Lock L3 & rgb[2]	L_3	-	[0, 1]	0.40
G[4]	Key K1	K_1	-	[0, 1]	0.10
G[5]	Key K2	K_2	-	[0, 1]	0.30
G[6]	Key K3	K_3	-	[0, 1]	0.10
G[7]*	Maximum age	a_{max}	s	[20, 300]	200
G[8]*	Initial Growth: Free Energy ratio	g_0	-	[0.2, 0.8]	0.5
G[9]*	Maximum Force/Kg	F_{max}	-	[0, 35]	30
G[10]	-	-	-	-	-
G[11]	-	-	-	-	-
G[12]	Reward for currency C1	θ_1	-	[-1, 1]	0.10
G[13]	Reward for currency C2	θ_2	-	[-1, 1]	0.10
G[14]	Reward for currency C3	θ_3	-	[-1, 1]	0.10
G[15]	Reward for currency C4	θ_4	-	[-1, 1]	0.30
G[16]	Tag number	-	-	≥ 1	1
G[17]*	Reproduction min. required mass fraction	m_{repMin}	-	[0.2, 1]	0.75
G[18]	Offspring per reproduction	$n_{offspring}$	-	≥ 1	1
G[19]*	Relative mass of offspring	$o_{relMass}$	-	[0, 0.5]	0.2
G[20]	-	-	-	-	-
G[21]	Reward for consumption	θ_5	-	[0, 1]	0.25
G[22]	Minimum viable body mass fraction	f_{starv}	-	[0.2 - 0.9]	0.75
G[23]	Time for single rotation	t_{rot}	s	[0.9 - 1.1]	1
G[24]	Digestion rate constant	ξ_{con}	s^{-1}	[0, 1]	0.2
G[25]	Max change in Growth: Free Energy	$g_{\Delta max}$	-	≥ 0	0
G[26]	Reward for reproduction	θ_6	-	[0, 1]	1
G[27]*	Mean reproductions per max lifetime	λ_{rMax}	-	≥ 1	4
G[28]	Alpha for currency C1	α_{C1}	-	≥ 0	1.87e-3
G[29]	Alpha for currency C2	α_{C2}	-	≥ 0	1.25e-2
G[30]	Alpha for currency C3	α_{C3}	-	≥ 0	1.83e-2
G[31]	Alpha for currency C4	α_{C4}	-	≥ 0	1.13e-3
G[32]	Alpha for consumption	α_{eat}	-	≥ 0	6.2e-1
G[33]	Alpha for reproduction	α_{rep}	-	≥ 0	8.29e-1
G[34]*	Energy reserve (C0)	C_{0res}	-	≥ 0	1
G[35]	Synthesizability of C1	z_1	-	{0, 1}	1
G[36]	Synthesizability of C2	z_2	-	{0, 1}	0
G[37]	Synthesizability of C3	z_3	-	{0, 1}	1
G[38]	Synthesizability of C4	z_4	-	{0, 1}	0
G[39]	-	-	-	-	-

Table A.4: Ent Gene values 0-39. * indicates the gene values which change under evolution.

Gene	Description	Symbol	Units	Range	Initial Value
G[40]	Entity type	-	-	≥ 1	1
G[41]*	Sensing radius	$r_{sensing}$	m	≥ 0	3
G[42]*	Maximum gene difference	$\Delta_{g_{max}}$	-	$[0, 0.2]$	0.15
G[43]*	Initial currency provided	$C_{initial}$	-	≥ 0	0.1
G[44]	Impact damage coefficient	k	-	≥ 0	0.001
G[45]	Consumption conversion efficiency	ξ_{con}	-	≥ 0	0.9
G[46]*	Maximum velocity	v_{max}	m/s	$[0, 3]$	1
G[47]	Self mass conversion efficiency	ψ	-	≥ 0	0.9
G[48]	Movement cost rate	ν_k	-	≤ 0	-4e-5
G[49]	Transaction cost	k_{trans}	-	≥ 0	0
G[50]	Currency amount given	C_{given}	-	≥ 0	0.1
G[51]	Base metabolic running cost	k_{base}	E/kg	≤ 0	-0.002
G[52]*	Synthesized currency	C_{synth}	-	0 - 1	0.02
G[53]	Pain reward	θ_{pain}	-	$[-1, 1]$	-1
G[54]	Number of Locks	n_{locks}	-	≥ 1	3
G[55]	Energy density	ρ_{Ent}	E/kg	≥ 0	1
G[56]*	Gene copying noise	$\delta_{G[i]}$	-	$[0.01, 0.05]$	0.015
G[57]*	Stomach mass ratio	q_f	-	$[0.05, 0.5]$	0.2
G[58]	C0 given to offspring at birth	$C_{0,0}$	-	≥ 0	1
G[59]	-	-	-	-	-
G[60]	Beta value for currency C1	β_{C1}	-	≥ 0	3.8e-3
G[61]	Beta value for currency C2	β_{C2}	-	≥ 0	3.8e-3
G[62]	Beta value for currency C3	β_{C3}	-	≥ 0	3.8e-3
G[63]	Beta value for currency C4	β_{C4}	-	≥ 0	1.14e-2
G[64]	Beta value for currency consumption	β_{eat}	-	≥ 0	9.5e-3
G[65]	Beta value for currency reproduce	β_{rep}	-	≥ 0	5e-3
G[66]*	Currency 1 amount given	C_{1given}	-	≥ 0	0.1
G[67]*	Currency 2 amount given	C_{2given}	-	≥ 0	0.1
G[68]*	Currency 3 amount given	C_{3given}	-	≥ 0	0.1
G[69]	Currency 4 amount given	C_{4given}	-	≥ 0	1

Table A.5: Ent Gene values 40-69. * indicates the gene values which change under evolution.

A.3 Repeat runs

The stochastic nature of the simulated environment and of the learnt policy leads to variation between repeated runs of the same experimental conditions. The figures below provide data for repeat runs of the main experiments (see also Table 3 for details). The data shown in figures below are: Ent population (first column), the total number of coins (or vitamins/poisons) being collected per successive 50 s period (column 2); the ratio of these two, showing mean number of coins collected per 50 s increment per Ent (column 3); reward coefficient values for eating, θ_3 (column 4); reward coefficient values for coin collection, θ_4 (column 5). Two styles of plots are used: (i) Where the majority of populations survive the experiment (see values in square brackets in each caption), median values for the surviving populations are shown with a blue band indicating the full range of those cases. A locally weighted linear regression method (the ‘Lowess’ Matlab function, with a window of 500 points). For comparison, an example raw (non-smoothed) population time series is shown in each case. (ii) Where populations do not survive the experiment, data for each population are shown as individual time series (in this case only the reward coefficient time series are smoothed.). In the captions below, results presented are linked to experiment numbers and coin increase rates defined in Table 2 (*faster* rate of coin increase being $0.014s^{-1}$ and the *slower* $0.0097s^{-1}$).

A.3.1 COINS EXPERIMENTS

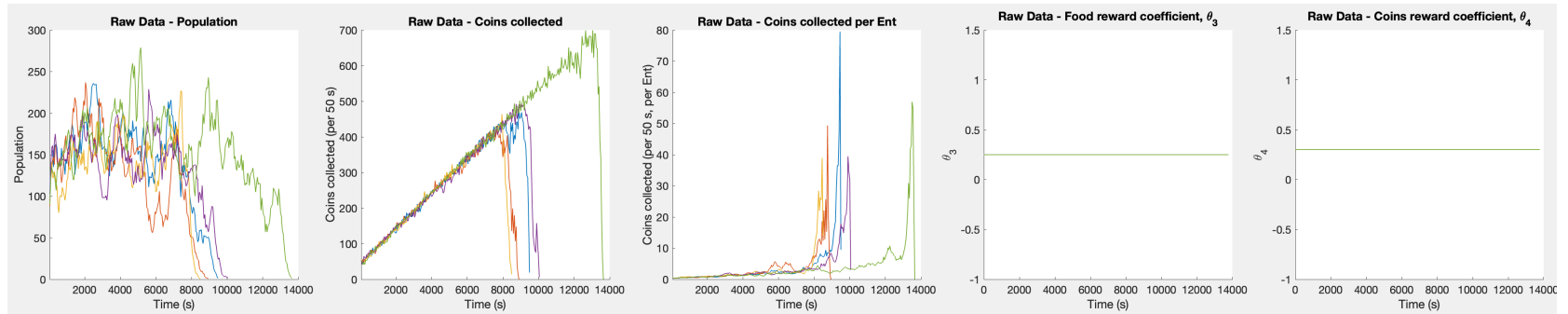


Figure 9: Experiment 2c: Learning, slower coin increase.

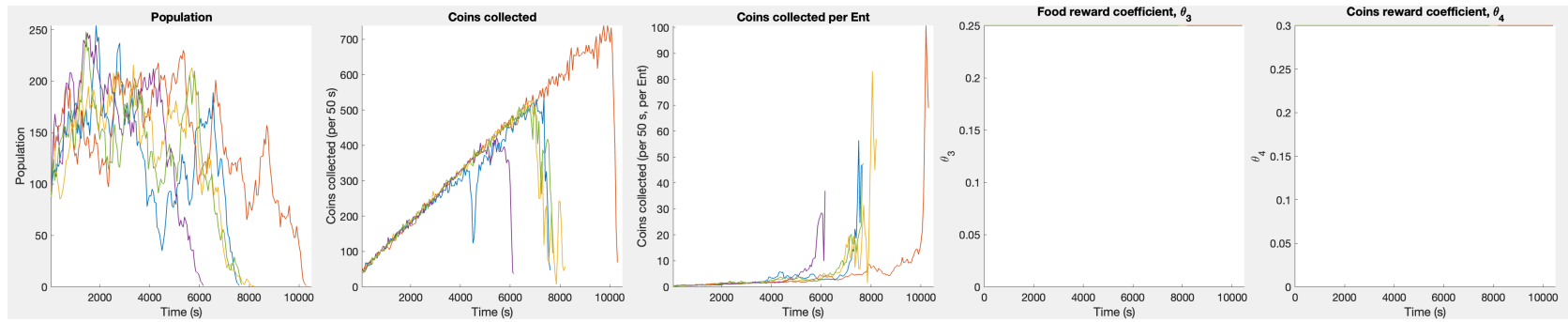


Figure 10: Experiment 2c: Learning, faster coin increase.

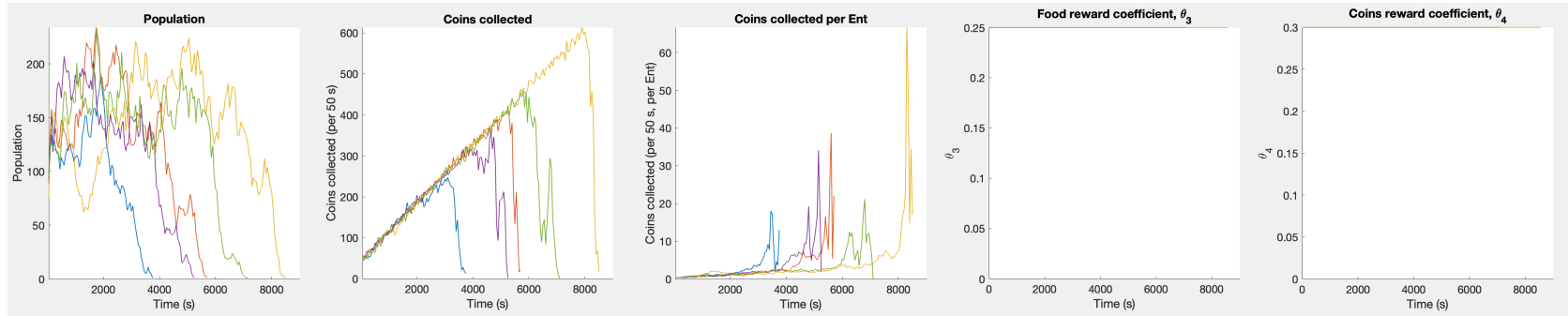


Figure 11: Experiment 2d: Learning and Evolution, faster coin increase.

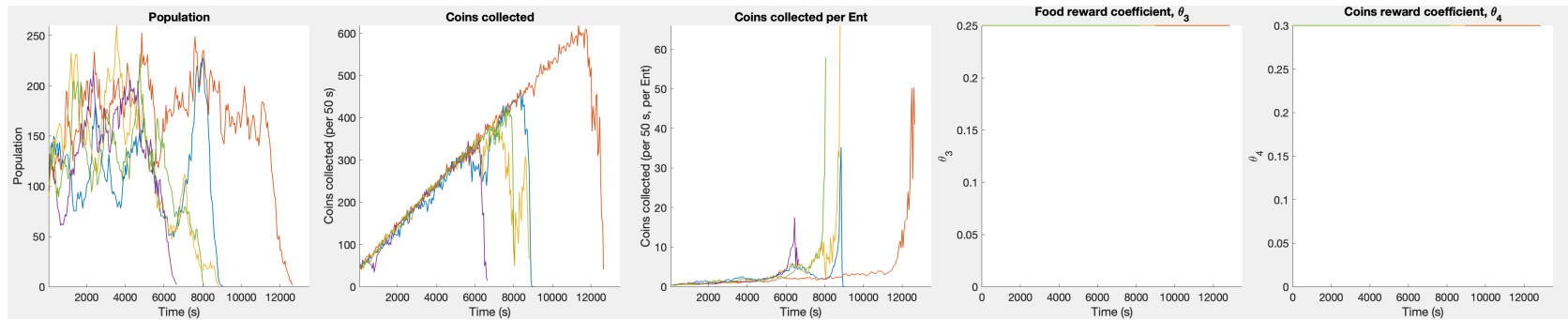


Figure 12: Experiment 2d: Learning and Evolution, slower coin increase.

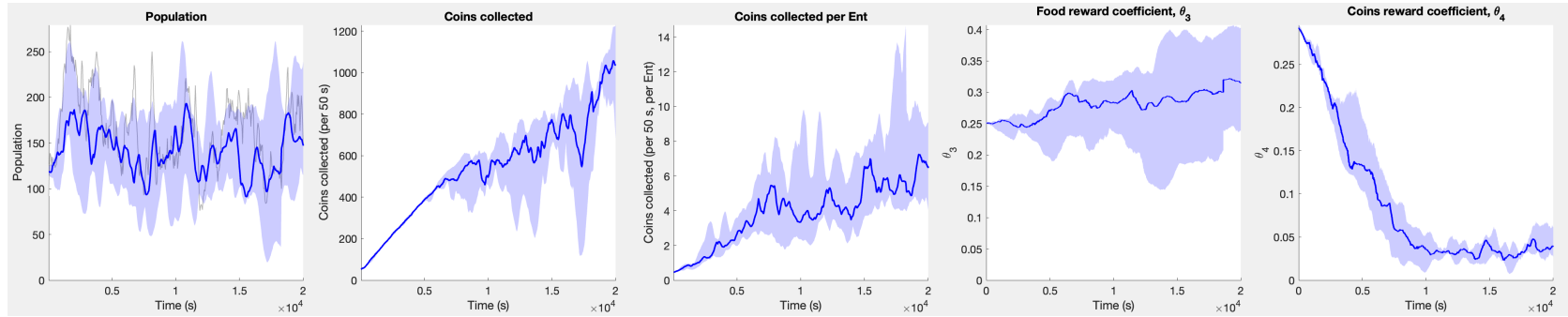


Figure 13: Experiment 2e: Learning and Reward Updating (RULE), faster coin increase.

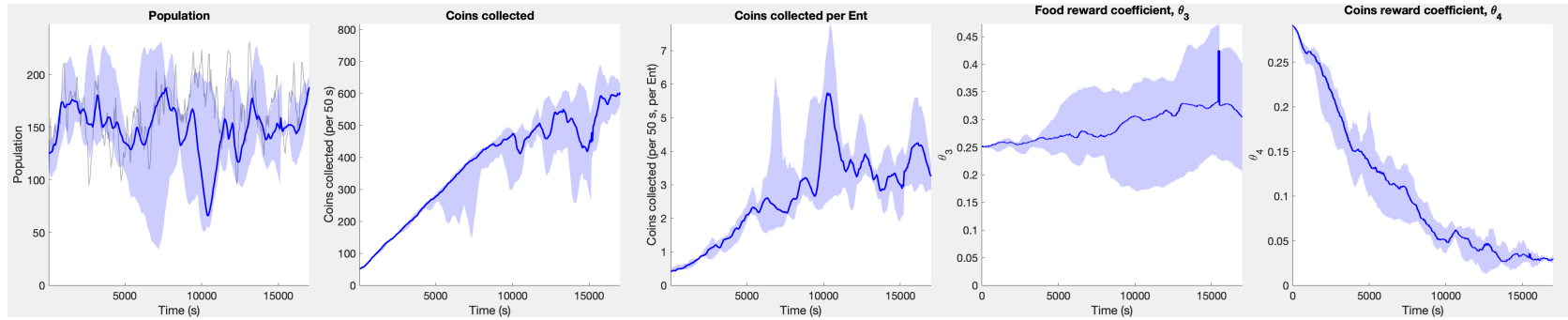


Figure 14: Experiment 2e: Learning and Reward Updating (RULE), slower coin increase.

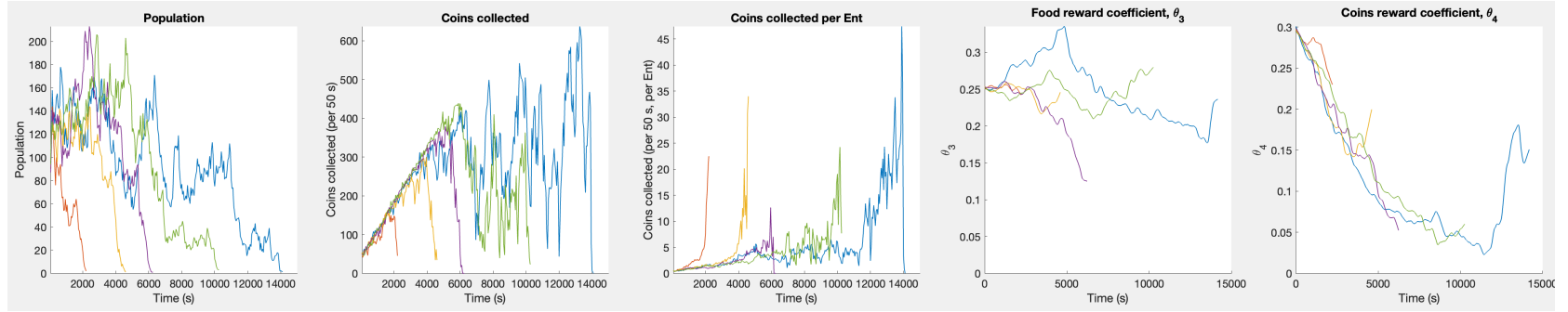


Figure 15: Experiment 2f: Learning, Reward Updating (RULE) and Evolution, faster coin increase.

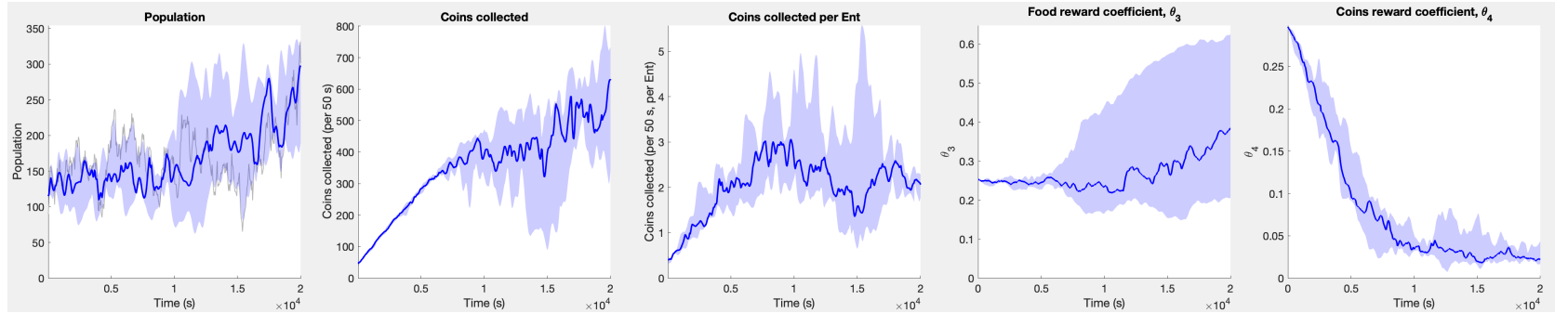


Figure 16: Experiment 2f: Learning, Reward Updating (RULE) and Evolution, slower coin increase.

A.3.2 VITAMINS AND POISONS EXPERIMENTS

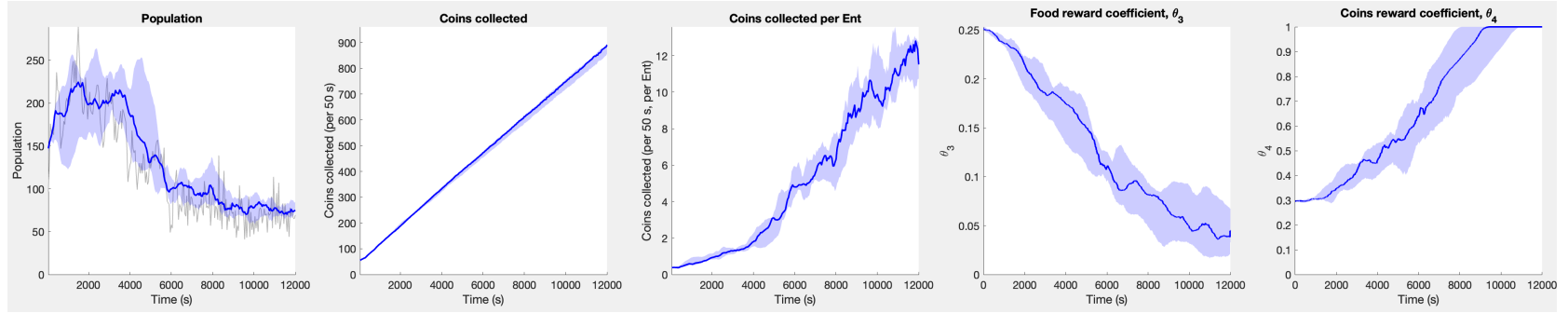


Figure 17: Experiment 3: Weaker vitamins, Learning, Reward Updating (RULE), faster coin (vitamin) increase.

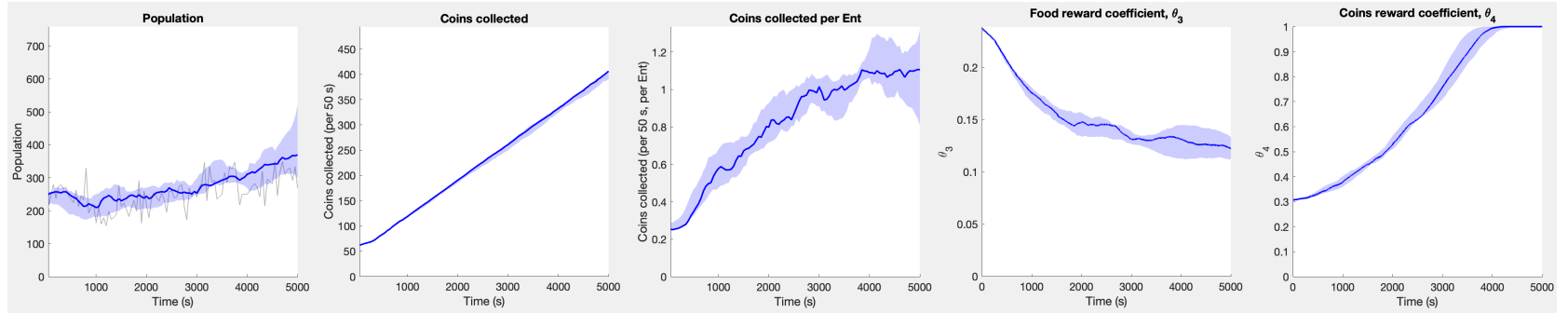


Figure 18: Experiment 3: Stronger vitamins, Learning, Reward Updating (RULE), faster coin (vitamin) increase.

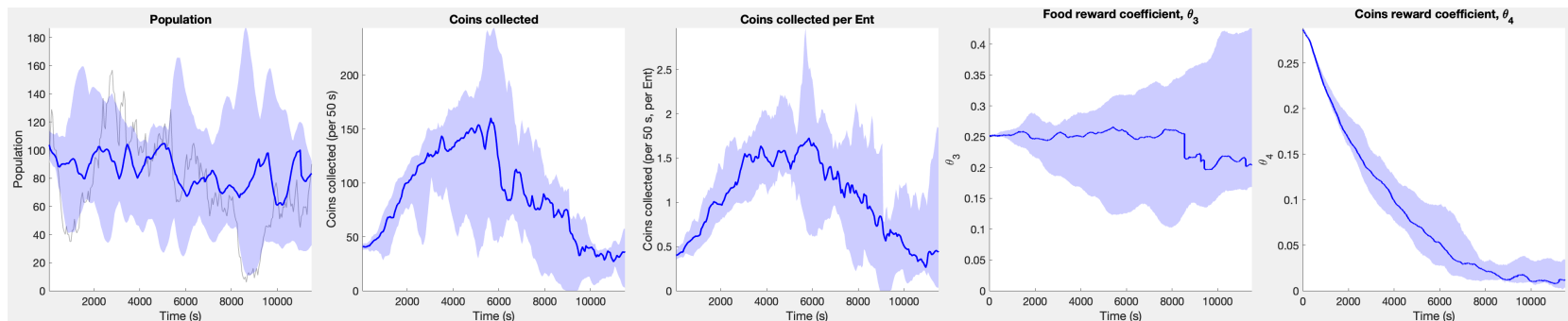


Figure 19: Experiment 3: Weaker poison, Learning, Reward Updating (RULE), slower coin (poison) increase.

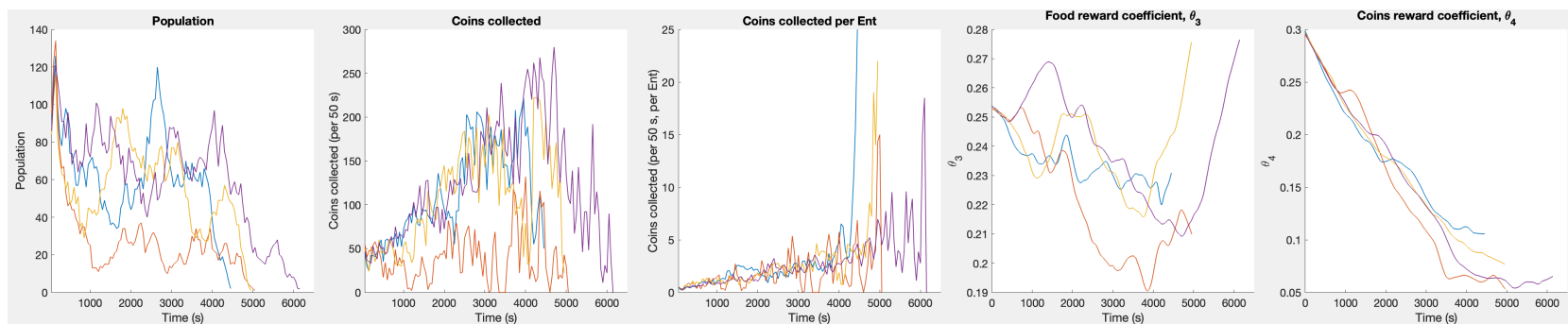


Figure 20: Experiment 3: Weaker poison, Learning, Reward Updating (RULE), faster coin (poison) increase.

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