# Genetic Team Composition and Level of Selection in the Evolution of Cooperation

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Abstract—In cooperative multiagent systems, agents interact to solve tasks. Global dynamics of multiagent teams result from local agent interactions, and are complex and difficult to predict. Evolutionary computation has proven a promising approach to the design of such teams. The majority of current studies use teams composed of agents with identical control rules ("genetically homogeneous teams") and select behavior at the team level ("team-level selection"). Here we extend current approaches to include four combinations of genetic team composition and level of selection. We compare the performance of genetically homogeneous teams evolved with individual-level selection, genetically homogeneous teams evolved with team-level selection, genetically heterogeneous teams evolved with individual-level selection, and genetically heterogeneous teams evolved with team-level selection. We use a simulated foraging task to show that the optimal combination depends on the amount of cooperation required by the task. Accordingly, we distinguish between three types of cooperative tasks and suggest guidelines for the optimal choice of genetic team composition and level of selection.

*Index Terms*—Altruism, artificial evolution, cooperation, evolutionary robotics, fitness allocation, multiagent systems (MAS), team composition.

# I. INTRODUCTION

**M** ULTIAGENT SYSTEMS (MAS) span a large number of research fields, from software agents to robotics, and play a key role in several industrial applications, such as ground and air vehicle control, supply chains, or network routing. The design of control rules for MAS is challenging because agent behavior depends not only on interactions with the environment but also on the behavior of other agents. As the number of interacting agents in a team grows, or when agent behaviors become more sophisticated, the design of suitable control rules rapidly becomes very complex. This is especially true when agents are expected to coordinate or cooperate to collectively achieve a desired task. Evolutionary computation has been advocated as an effective and promising strategy to generate control parameters and decision rules for collective agents [1], [2].

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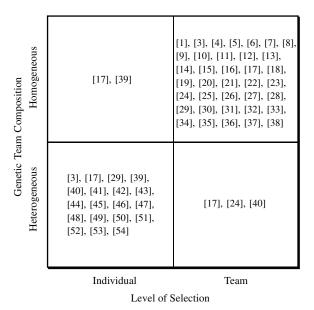


Fig. 1. Sample of approaches to the evolution of multiagent teams. The majority of work uses genetically homogeneous teams, usually created from a cloned individual genome, with team selection. In some cases, authors created behaviorally heterogeneous agents out of a single team genome: Luke [25], [26] decoded team genomes into six separate sub-teams with one or two identical players each. Other authors [11], [20], [22], [28], [33] decoded one team genome into different single agent genomes. Yet another approach was taken by work using distributed embodied evolution to evolve heterogeneous teams [43], [50]-[54]. In these cases, selection and replication were entirely distributed among agents, with dynamics reminiscent of the replicator dynamics observed in bacterial evolution [55] and game theoretic models [56]. In some cases, teams were evolved using a continuously updated gene-pool rather than separate gene-pools for subsequent generations ("steady state evolution") [45]-[47]. Finally, some authors have conducted more detailed comparisons of the influence of genetic team composition or level of selection alone: Martinoli [40] also considered more complex methods of selection. Stanley et al. [49] clustered genetically similar individuals into sub-teams that shared fitness, which resulted in partially heterogeneous teams. Mirolli et al. [39] also compared partially heterogeneous teams. Quinn [29] evaluated individuals in different heterogeneous teams to create robust homogeneous teams.

In addition to the methodological issues of evolving agents that operate in isolation [2], the evolution of agent teams must address two major issues: 1) It must determine optimal team composition. Agents of a team may either share control rules (genetically homogeneous teams) or employ different ones (genetically heterogeneous teams), and 2) It requires a suitable method for selective reproduction of desired team behavior. Selection may operate either on individuals (individual-level selection) or on teams (team-level selection). In the simplest case, one must decide between genetically homogeneous or

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heterogeneous teams, and between selecting agents at the individual or at the team level.

Fig. 1 shows a sample of previous work on the evolution of MAS in robotics, combinatorial optimization, cellular automata, artificial life, genetic programming, and others, plotted according to the chosen genetic team composition and level of selection. In addition to work cited in Fig. 1, some authors have used cooperative co-evolutionary algorithms (CCEAs, [57]) to evolve heterogeneous control rules for teams of agents [58]-[61]. CCEAs are applied by decomposing problem representations into subcomponents and then creating a separate population of individuals for each subcomponent. This approach allows teams to be composed of specialized subgroups and corresponds to the biological co-evolution of multiple species. In their basic form, CCEAs require the designer to manually decompose the multiagent task, and thus to solve part of the optimization problem beforehand. Work that used machine learning techniques other than evolutionary computation (e.g., reinforcement learning) is not considered in this paper.

Fig. 1 suggests that the majority of current approaches to the evolution of MAS use genetically homogeneous teams evolved with team-level selection. Where the reasons for the choice of genetically homogeneous teams are made explicit, it is argued that homogeneous teams are easy to use [8], [36], require fewer evaluations [25], [32], scale more easily [13], and are more robust against the failure of team members [13], [62] than heterogeneous teams. Many other approaches use genetically heterogeneous teams evolved with individual-level selection. Genetically heterogeneous teams are sometimes seen as providing more behavioral flexibility [25] and as providing advantages in tasks that require specialization [7], [25], [62], [63].

The terms "homogeneous team" and "heterogeneous team" used in the current literature cover many different aspects. It is important to note that while all agents in genetically homogeneous teams share the same genes, agents can nevertheless be behaviorally heterogeneous. This can happen when agents differentiate during their lifetime, for example due to varying initial conditions [30], or due to developmental processes or learning [64]. This can also happen when agents "activate" different parts of their genome, for example when each agent's behavior is controlled by a different section of a single team genome [11], [22], [28], [33]. In this case, agents can specialize on different functions, yet be genetically identical, just like specialized cells in a biological organism. Conversely, it is important to note that genetically heterogeneous teams are those in which agents are, on average, not genetically more similar to team members than to agents in the rest of the population [65], [66]. This means that teams resulting from embodied evolution or common versions of steadystate evolution are usually genetically heterogeneous although these algorithms often generate multiple offspring from a single parent, resulting in genetically similar (but not identical) agents. In some cases, teams consist of clonal sub-teams [25], [26] or agents that share only part of their genome. Teams with agents that are, on average, genetically more similar (but not identical) to members of their team than to members of the rest of the population are termed "partially heterogeneous." The effects of partial genetic heterogeneity on the evolution of multiagent teams are not yet fully explored in evolutionary computation [39], but there is evidence that they can lead to improved specialization [25], [26]. These effects have been deeply studied in biology [67], [68].

The choice of level of selection is rarely discussed explicitly. Some research has addressed the related issue of credit assignment for the evolution of MAS [40], [69]. In the context of MAS, credit assignment is concerned with distributing fitness rewards among individual agents. Fitness distribution leads to credit assignment problems [70], [71] in many cooperative multiagent tasks, because individual contributions to team performance are often difficult to estimate or difficult to monitor [72]. Selection is usually performed on the basis of accumulated individual or team fitness, which may be the result of many fitness rewards with different types of credit assignment. Therefore an optimal choice of level of selection is not only influenced by the type of task but also by the types of credit assignment used.

Genetic team composition and level of selection have long been identified as two important factors for the evolution of biological multiagent teams such as groups of genes, cells, individuals, or other replicators [67], [73]. In particular the evolution of altruism [74], in which agents cooperate to increase team fitness in spite of an individual fitness cost to the cooperator, has received a lot of attention [68], [75]. Here we define cooperation as a behavior that increases the fitness of other agents, and altruistic cooperation (altruism) as a behavior that increases the fitness.

In this paper, we focus on cooperative multiagent tasks that do not require specialization. We compare the performance of robot teams evolved in four evolutionary conditions: genetically homogeneous teams evolved with team-level selection; genetically homogeneous teams evolved with individual-level selection; genetically heterogeneous teams evolved with teamlevel selection; and genetically heterogeneous teams evolved with individual-level selection. We evaluate the performance of robot teams evolved in these four evolutionary conditions for three classes of multirobot tasks: a task that does not require cooperation; a task that requires cooperation but does not imply a cost for cooperators; and a task that requires altruistic cooperation, i.e., a task that implies an individual fitness cost for cooperators. Cooperative tasks that benefit from specialization were not considered in this paper.

#### **II. EVOLUTIONARY CONDITIONS**

The four possible combinations of genetic team composition and level of selection were formalized into four evolutionary algorithms (Fig. 2). For the remainder of the paper we will use the terms "homogeneous" and "heterogeneous" to designate genetically homogeneous and genetically heterogeneous teams, respectively, and the terms "individual selection" and "team selection" to designate teams evolved with individuallevel selection and team-level selection, respectively. We consider populations composed of M teams, each composed of N

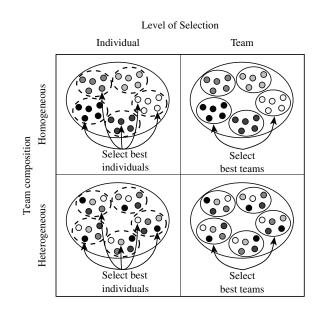


Fig. 2. Four evolutionary conditions. A population (large oval) is composed of several teams (medium ovals), each of which is composed of several robots (small circles) evaluated together. Genetic team composition is varied by either composing teams of robots with *identical genomes* (homogeneous, identical shading), or *different genomes* (heterogeneous, different shading). Level of selection is varied by selecting *teams* (team selection), or selecting *individuals*, independent of their team affiliation (individual selection).

Algorithm 1 Homogeneous teams, individual selection
for each of $M$ new teams do
select two individuals from all old teams
recombine their genomes to create one new genome
mutate new genome
clone new genome to obtain $N$ genomes for new team
end for

individuals. Population size and team sizes are kept constant across generations. At each generation, the old population is entirely replaced by a new population of offspring. Individuals' genomes are binary strings.

1) Algorithm 1—Homogeneous Teams, Individual Selection: Each of the M teams at generation 0 was formed by generating one random genome and cloning it N-1 times to obtain N identical robot genomes (clones) per team. Teams were evaluated in the task and an individual fitness determined for each of the N robots. For a new generation, each of the Mnew teams was created from two *individuals* selected among all individuals of all old teams in the population using roulette wheel selection. The two genomes of the selected individuals were recombined (one-point crossover, crossover probability of 0.05) to produce one new genome. The resulting new genome was mutated by flipping the value of each bit with a probability of 0.05 and then cloned N-1 times to generate the N robot genomes of the new team. Teams evolved using this evolutionary condition were thus genetically homogeneous.

2) Algorithm 2—Homogeneous Teams, Team Selection: Each of the M teams at generation 0 was formed by generating one random genome and cloning it N - 1 times to obtain N identical robot genomes (clones) per team. Teams were

Algorithm 2 Homogeneous teams, team selection	
for each of $M$ new teams do	
select two old teams	
recombine their genomes to create one new ge	enome
mutate new genome	
clone new genome to obtain $N$ genomes for n	iew team
end for	
Algorithm 3 Heterogeneous teams, individual select	ction
for each of $M$ new teams do	
for each of $N$ new team members do	
select two individuals from all old teams	
recombine their genomes to create one new	genome

mutate new genome add new genome to new team

end for

end for

evaluated in the task, and for each team a team fitness was determined as the sum of the individual fitnesses of all N robots. For a new generation, each of the M new teams was created from two old *teams* selected using roulette-wheel selection. The two genomes of the selected teams were recombined (one-point crossover, crossover probability of 0.05) to produce one new genome. The resulting new genome was mutated by flipping the value of each bit with a probability of 0.05 and then cloned N - 1 times to obtain the N robot genomes of the new team. Teams evolved using this evolutionary condition were thus genetically *homogeneous*.

3) Algorithm 3—Heterogeneous Teams, Individual Selec*tion:* Each of the M teams at generation 0 was formed by generating N random genomes. Teams were evaluated in the task and an individual fitness determined for each of the N robots. For a new generation, each of the  $N \times M$  new individuals was created from two individuals selected among all individuals of all old teams in the population using roulette wheel selection. The two genomes of the selected individuals were recombined (one-point crossover, crossover probability of 0.05) to produce one new genome. The resulting new genome was mutated by flipping the value of each bit with a probability of 0.05. This process was repeated  $N \times M - 1$ times to form M new teams of N individuals each. In this evolutionary condition robots were not, on average, genetically more similar to team members than to robots in the rest of the population, and thus teams were genetically heterogeneous.

4) Algorithm 4—Heterogeneous Teams, Team Selection: Each of the M teams at generation 0 was formed by generating N random genomes. Teams were evaluated in the task, and for each team a team fitness was determined as the sum of the individual fitnesses of all N robots. For a new generation, each of the  $N \times M$  individuals was created from two old *teams* selected using roulette wheel selection. Two genomes, each randomly selected among the members of a selected team, were recombined (one-point crossover, crossover probability of 0.05) to produce one new genome. The resulting new genome was mutated by flipping the value of each bit with

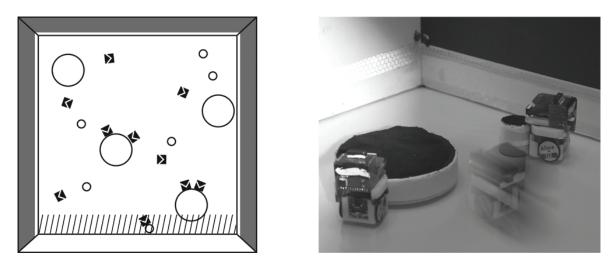


Fig. 3. Left: Experimental setup for task 3, the altruistic cooperative foraging task. Ten micro-robots (black squares with arrows) searched for small and large tokens and transported them to the target area (hatched area at bottom) under the white wall (the other three walls were black). An identical setup was used in the other two tasks, except that the arena contained either only small tokens in task 1, or only large tokens in task 2. Right: Three micro-robots in task 3, which is the altruistic cooperative foraging task. The robot in the background could transport the small token by itself. The robot at the left could not transport the large token by itself and needed to wait for the arrival of a second robot (blurred in the picture due to its rapid movement towards the large token).

Algorithm 4 Heterogeneous teams, team selection
for each of $M$ new teams do
for each of $N$ new team members do
select two old teams
randomly select two old team members
recombine their genomes to create one new genome
mutate new genome
add new genome to new team
end for
end for

a probability of 0.05. This process was repeated  $N \times M - 1$  times to form M new teams of N individuals each. In this evolutionary condition, robots were not, on average, genetically more similar to team members than to robots in the rest of the population, and thus teams were genetically *heterogeneous*.

#### III. EXPERIMENTAL METHOD

# A. Scenario

The experimental setup (Fig. 3) consisted of a  $50 \times 50$  cm<sup>2</sup> arena with 10 micro-robots and two types of tokens: small and large. We chose to study a foraging task, because foraging combines several aspects of multiagent tasks (distributed search, coordinated movement, transportation) and relates to many real-world problems [76], [77]. In addition, foraging is a wide-spread and well-studied behavior of many biological societies [78]–[80]. Experiments were conducted in simulation of micro-robots and evolved controllers were transferred to the real robots (see Section III-D).

Robots foraged tokens by transporting them into a 4-cm wide region at one side of the arena marked by a white wall. A single robot was sufficient to transport a small token. At least two robots were required to transport a large token, and thus retrieval of large tokens required cooperation. Cooperating

agents had to coordinate their behaviors to successfully align their positions before and during token transport.

The micro-robots [81] were small  $(2 \times 2 \times 4 \text{ cm}^3)$  twowheeled robots equipped with three infrared distance sensors at the front and one at the back, which could sense objects up to 3 cm away and allowed robots to distinguish between small and large tokens (Fig. 4 left). An extension module with a fourth infrared distance sensor with a range of up to 6 cm and a linear camera were mounted higher on the robot, overlooking tokens but sensitive to other robots and walls.

#### B. Control and Genetic Architecture

Robots were controlled by a feed-forward neural network with a single layer of three hidden neurons (Fig. 4 right) and a sigmoid activation function (tanh). The inputs were given by the activation values of five infrared sensors, two vision sensors, and a constant bias value of -1. Infrared sensor activation values were scaled in the range [0; 1]. Vision sensors were an average of three equidistally spread camera pixels spanning a field of view of 18°, for the left or right side of the image, respectively. The averages were thresholded to yield 0 for a white or 1 for a black arena wall. Using the average value of three pixels rather than a single pixel allowed a robust detection of the white foraging target area in spite of the presence of other robots in the field of view. The two output units were used to control the left and right wheel motors. The activation values in the range [-1; 1] were mapped into speeds in the range [-4; 4] cm/s, with speeds in the interval of [-2.5; 2.5] cm/s set to 0 because of unreliable motor response at low speeds.

The neural network connection weights were in the range of [-2; 2] and coded on 8 bits. The genome of one individual was thus  $8 \times 32$  bits long.

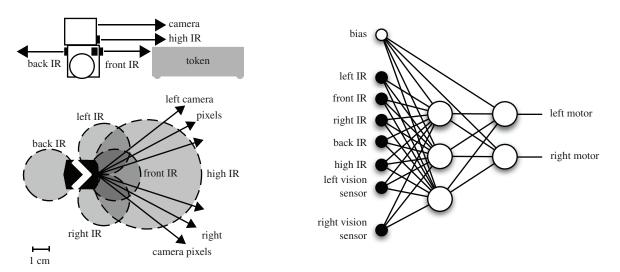


Fig. 4. Left: Side and top-view schematics of a simulated micro-robot. The robot was equipped with four infrared (IR) distance sensors (three at the front, one at the back) to detect tokens, and a camera to identify the target area. A fifth infrared distance sensor (high IR) was mounted higher on the robot and thus overlooked tokens. This allowed robots to distinguish tokens from walls and other robots. Right: The neural network architecture, which is a feed-forward neural network with a single layer of three hidden neurons. Inputs were given by the activation values of five infrared (IR) sensors and two vision sensors with activation values computed from left and right camera pixels (see text).

### C. Collective Tasks

We devised three types of foraging tasks that differed in the amount of cooperation required from agents.

1) Task 1—Individual Foraging: The arena contained 6 small tokens, each of which awarded 1 fitness point to the foraging robot. This task did not require cooperation, because a single agent was sufficient to transport a small token.

2) Task 2—Cooperative Foraging: The arena contained four large tokens, which each awarded 1 fitness point to each team member, irrespective of its participation in the token foraging. This corresponded to a situation where the individual contributions to team performance were not known, i.e., a situation with credit assignment problems [70], [71], which is the case for many cooperative multiagent tasks [72]. This task required cooperation because it could not be accomplished by a single agent.

3) Task 3—Altruistic Cooperative Foraging: The arena contained six small and four large tokens. Small tokens each awarded one fitness point to the foraging robot and large tokens each awarded one fitness point to each team member, irrespective of their participation in the token foraging. In this task cooperation was costly for individuals, because individuals that did not cooperate always had higher fitness than their cooperating team mates. This meant that cooperators suffered a relative individual fitness cost and therefore this task required altruistic cooperation [68].

#### D. Evolutionary Experiments

Due to the large number of evaluations required for the evolution of robot behaviors, all evolutionary experiments were conducted using a physics-based 2-D simulator [82], which is available as part of an open evolutionary framework [83]. All simulation parameters, including robot size, shape, speed, and weight, as well as collision dynamics, friction

forces, and sensor and actuator modalities, were based on the micro-robots described in Section III-A.

We evolved teams of robots under the four evolutionary conditions separately for each of the three tasks, making a total of 12 experimental lines. Evolutionary experiments lasted for 300 generations. Twenty independent runs were performed for each experimental line. Populations consisted of 100 teams of 10 agents each. Each team was evaluated 10 times for 3 minutes with random token and robot starting positions and orientations. Fitness was averaged over the 10 evaluations.

To compare the efficiency of the four evolutionary conditions, we re-evaluated the best teams at generation 300 for 1000 times and compared their team fitness. Fitness values were analyzed using Wilcoxon rank sum tests. All fitness values were normalized for each task, with 0 being the minimal possible fitness and 1 the theoretical maximum value.

#### **IV. RESULTS**

#### A. Task 1—Individual Foraging

Successful foraging behavior evolved for all four evolutionary conditions (Fig. 5). After 300 generations of artificial evolution, heterogeneous teams evolved with individual selection collected all 10 tokens in most evaluations and achieved fitness values close to the maximum value achievable. These fitness values were higher than those of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection (Wilcoxon rank sum test, df = 38, P < 0.001 and P < 0.006, respectively). A possible reason could be the disparities in genome evaluation in homogeneous and heterogeneous teams. For a team size of N agents, heterogeneous teams evaluated N times more genomes than homogeneous teams. This was because each heterogeneous team consisted of N different genomes, whereas homogeneous teams consisted of N identical genomes. On the other hand, homogeneous teams evaluated each genome N times more

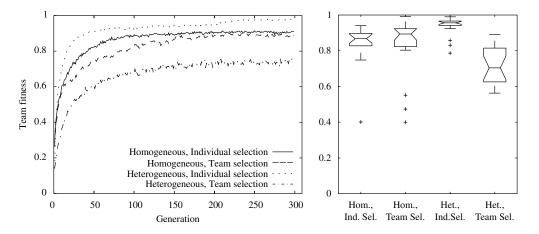


Fig. 5. Task 1—Individual Foraging. Left: Evolution of the best team fitness averaged over the best teams in 20 independent evolutionary runs over 300 generations. Right: The best team at generation 300 of each of the 20 independent experiments per evolutionary condition and per task was evaluated 1000 times. The mid-line in the box is the median, while the box represents the upper and lower quartile above and below the median. The bars outside the box generally represent the maximum and minimum values, except when there are outliers, which are shown as crosses. We define outliers as data points which differ more than 1.5 times the interquartile range from the border of the box. The notches represent the uncertainty in the difference of the medians for box-to-box comparison. Boxes whose notches do not overlap indicate that the medians differ at the 5% significance level [84]. In this task, which did not generation, heterogeneous teams evolved with individual selection performed best, followed by homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection. Heterogeneous teams evolved with team selection performed significantly worse than all other evolutionary conditions.

often than heterogeneous teams. This was because each team evaluation evaluated an identical genome N times. Our results suggest that higher evaluation accuracy may have been less important than a larger number of different genomes in this task. The larger number of genomes may have allowed heterogeneous teams to discover solutions faster than homogeneous teams, which could explain the steep initial fitness increase. It may also have allowed heterogeneous teams, which could explain the steep initial fitness increase. It may also have allowed heterogeneous teams to discover better solutions than homogeneous teams, which could explain the higher final fitness obtained with this evolutionary condition. To test whether these disparities in genome evaluation caused the high team performance of heterogeneous teams evolved with individual selection, we performed a set of additional experiments (see additional experiments without these disparities in the next section).

Performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection did not differ significantly (P = 0.337). This was because with roulette-wheel selection the probability of a team to be selected was the same as the sum of the probabilities of each individual team member to be selected. Since all team members of homogeneous teams shared the same genome, selection probabilities for a given genome were equal for both homogeneous evolutionary conditions. It should be noted, however, that this is not necessarily true for other types of selection. Selection mechanisms where the fitness of a genome is not directly proportional to its probability to be selected (e.g., truncation or rank-based selection) may lead to differences in the number of selected individuals with a given genotype and consequently affect the relative performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection. In these cases, individual selection may select for genomes that lead to higher maximum but lower average individual performance. However, additional investigations using truncation selection (selection of best 30% of the population; all other experimental

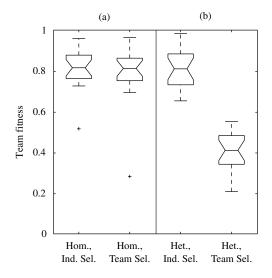


Fig. 6. Task 1—Individual foraging without disparities in genome evaluation. (a) Homogeneous teams evolved with one evaluation per team (instead of 10) and (b) heterogeneous teams evolved with 100 agents per population (instead of 1000). Heterogeneous teams evolved with individual selection performed similar to homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection in this task. For boxplot explanations, see Fig. 5.

parameters identical) did not find such performance differences in any of the three types of task (P = 0.350/0.903/0.394for tasks 1/2/3 respectively; see Fig. S4 in the supplementary material<sup>1</sup>).

Heterogeneous teams evolved with team selection performed significantly worse than all other evolutionary conditions (all three P < 0.002). This was because, unlike all other three evolutionary conditions, this evolutionary condition did not allow a direct link between the performance of a genome and its probability to be selected. Instead, selection

<sup>1</sup>An electronic supplement for this paper is available online at http://lis.epfl.ch/documentation.php

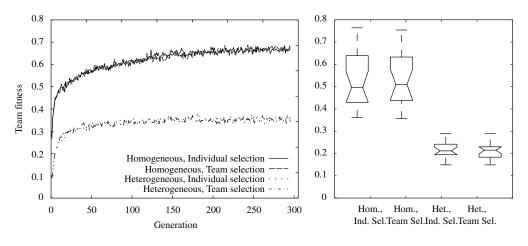


Fig. 7. Task 2 - Cooperative Foraging. Left: Evolution of the best team fitness averaged over the best teams in 20 independent evolutionary runs over 300 generations. Right: The best team at generation 300 of each of the 20 independent experiments per evolutionary condition and per task was evaluated 1000 times. Homogeneous teams performed significantly better than heterogeneous teams. For boxplot explanations, see Fig. 5.

of good genomes could only happen indirectly, by selecting those teams that contained a better mix of genomes than other teams. Since good genomes could be part of bad teams or bad genomes part of good teams, selection for good individual genomes was inefficient. This explains the slow initial fitness increase and the lowest final fitness of heterogeneous teams evolved with team selection.

# B. Task 1—Individual Foraging: Disparities in Genome Evaluation and Credit Assignment

To test the hypothesis that the high team performance of heterogeneous teams evolved with individual selection was caused by disparities in genome evaluation, we performed a set of additional experiments. First, we evolved homogeneous teams in the same task, but used only one evaluation per team rather than 10 evaluations [Fig. 6(a)]. Second, we evolved heterogeneous teams in the same task, but used only 100 agents per population rather than 1000 agents [Fig. 6(b)]. In this set of experiments homogeneous and heterogeneous teams therefore evaluated the same number of genomes and had the same number of evaluations per genome.

Without disparities in genome evaluation, heterogeneous teams evolved with individual selection performed similar to homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection (all three P > 0.597). Heterogeneous teams evolved with team selection performed worse than all other evolutionary conditions (all three P < 0.001), because the efficiency of selection was not affected by the changes in genome evaluation.

#### C. Task 2—Cooperative Foraging

Successful foraging behavior evolved for all four evolutionary conditions (Fig. 7). The experiments with a cooperative task led to a change in the relative performance of the four evolutionary conditions. Performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection was significantly higher than that of heterogeneous teams evolved with individual selection and heterogeneous teams evolved with team selection (all four P < 0.001), with the best fitness values in homogeneous teams up to 70% higher than those in heterogeneous teams. One possible reason is disparities in genome evaluation between homogeneous and heterogeneous teams (see next section). Another possible reason is that selection of good genomes could only happen indirectly in this task, which may have led to inefficient selection just as in heterogeneous teams evolved with team selection in task 1 (Section IV-B). This could be because fitness in this task was assigned to all team members, irrespective of their participation in the token foraging.

Performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection did not differ significantly (P = 0.839). This was because fitness in this task was assigned to all team members, irrespective of their participation in the token foraging. For the same reason, performance of heterogeneous teams evolved with individual selection and heterogeneous teams evolved with team selection did not differ significantly (P = 0.365).

# D. Task 2—Cooperative Foraging: Disparities in Genome Evaluation and Credit Assignment

To test the hypothesis that the differences in performance of heterogeneous teams that evolved with individual selection and homogeneous teams that evolved with individual selection and team selection were caused by disparities in genome evaluation or by the fitness assignment to all team members, we performed two sets of additional experiments. First, we again corrected for the disparities in genome evaluation. However, correcting for this factor alone did not eliminate the performance differences (see Fig. S2 in the supplementary material<sup>2</sup>). Second, we performed experiments where we again corrected for the disparities in genome evaluation and where fitness was only assigned to team members that participated in the token foraging. In these experiments, each of the four large tokens awarded 5 fitness points to each of the two transporting

<sup>2</sup>An electronic supplement for this paper is available online at http://lis.epfl.ch/documentation.php

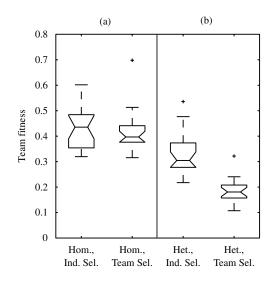


Fig. 8. Task 2—Cooperative foraging without disparities in genome evaluation and without credit assignment problems. (a) Homogeneous teams evolved with one evaluation per team (instead of 10) and (b) heterogeneous teams evolved with 100 agents per population (instead of 1000). The performance of heterogeneous teams evolved with individual selection was higher than the performance of heterogeneous teams evolved with team selection, but did not reach that of homogeneous teams in this task. For boxplot explanations, see Fig. 5.

robots, rather than 1 fitness point to each of the 10 team members. This second additional set of experiments therefore corresponded to a situation where the individual contributions to team performance were known, i.e., a situation without credit assignment problems.

Without the disparities in genome evaluation and without credit assignment problems, heterogeneous teams that evolved with individual selection outperformed heterogeneous teams that evolved with team selection (P < 0.001). This was because selection of good genomes could now happen directly, which allowed for efficient selection. However, the performance of heterogeneous teams evolved with individual selection remained lower than that of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection (P < 0.001 and P < 0.002, respectively, Fig. 8). A possible reason is that heterogeneous teams had to solve a more complex optimization task than homogeneous teams. Successful cooperation in heterogeneous teams required individuals to evolve behaviors to coordinate their actions with N-1 different team members, while individuals in homogeneous teams only had to evolve behaviors to coordinate with a single type of team member. In other words, homogeneous teams led to a smaller search space because all team members were per definition identical, and thus only a subset of the total number of possible team compositions was considered in these teams. Furthermore, individuals in heterogeneous teams were not just different in a team, but team members changed from one generation to the next. Both factors may have hindered the evolution of cooperative behavior in heterogeneous teams.

The performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection did not differ significantly (P = 0.441) in this second additional set of experiments. Heterogeneous teams evolved with team selection performed worse than all other evolutionary conditions due to inefficient selection (all three P < 0.001).

#### E. Task 3—Altruistic Cooperative Foraging

Successful foraging behaviors evolved for all four evolutionary conditions (Fig. 9). Team performance in the altruistic cooperative foraging task was systematically lower than in the cooperative foraging task. This may seem surprising because the larger number of tokens in the arena increased the total number of fitness points available. A possible reason is that the increased number of tokens led to more clutter in the arena, which made successful token transport more difficult (see video supplied with supplementary material<sup>3</sup>).

Homogeneous teams achieved significantly higher fitness values than heterogeneous teams (all four P < 0.001). Possible reasons are disparities in genome evaluation and inefficient selection for the foraging of large tokens because fitness points gained from large tokens were assigned to all team members, irrespective of their participation in the token foraging (see next section).

Performance of homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection did not differ significantly (P = 0.310). This was because selection probabilities for a given genome were again equal for both homogeneous evolutionary conditions.

looseness1Performance of heterogeneous teams evolved with individual selection and heterogeneous teams evolved with team selection did not differ significantly (P = 0.490). However, the four evolutionary conditions resulted in different foraging strategies in this task (Fig. 10): While homogeneous teams evolved with individual selection and homogeneous teams evolved with team selection as well as heterogeneous teams evolved with team selection collected a significantly higher proportion of large tokens than small tokens (all three P < 0.001), heterogeneous teams evolved with individual selection collected a significantly higher proportion of small tokens than large tokens (P < 0.001). In comparison to the other three evolutionary conditions, heterogeneous teams evolved with individual selection collected significantly high proportion of small tokens (all three P < 0.001), but significantly low proportion of large tokens of all four evolutionary conditions (all three P < 0.003).

# F. Task 3—Altruistic Cooperative Foraging: Disparities in Genome Evaluation and Credit Assignment

To test the hypothesis that the differences in performance of heterogeneous teams evolved with individual selection were caused by disparities in genome evaluation or by the fitness assignment to all team members, we performed two sets of additional experiments similar to those described in Section IV-D. First, we again corrected for the disparities in genome evaluation. However, correcting for this factor alone

<sup>&</sup>lt;sup>3</sup>An electronic supplement for this paper is available online at http://lis.epfl.ch/documentation.php

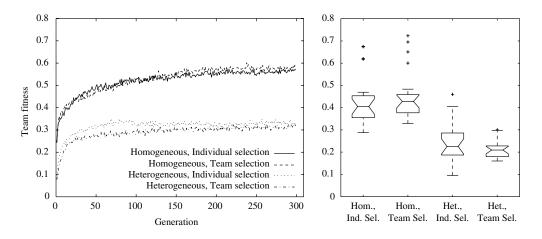


Fig. 9. Task 3—Altruistic Cooperative Foraging. Left: Evolution of the best team fitness averaged over the best teams in 20 independent evolutionary runs over 300 generations. Right: The best team at generation 300 of each of the 20 independent experiments per evolutionary condition and per task was evaluated 1000 times. Homogeneous teams performed significantly better than heterogeneous teams. For boxplot explanations, see Fig. 5.

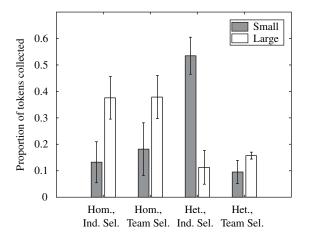


Fig. 10. Task 3—Altruistic cooperative foraging. The plot shows the average proportion of the six small tokens and four large tokens collected by the best teams at generation 300 for each of the 20 independent experiments and for each of the four evolutionary conditions. Heterogeneous teams evolved with individual selection pursued a different foraging strategy than teams of the other three evolutionary conditions, collecting very few large tokens but most small tokens.

did not eliminate the performance differences (see Fig. S3 in the supplementary material<sup>4</sup>). Second, we again performed experiments where we corrected for the disparities in genome evaluation and for credit assignment problems.

Without the disparities in genome evaluation and without credit assignment problems, heterogeneous teams evolved with individual selection outperformed heterogeneous teams evolved with team selection (P < 0.001). This was because selection of good genomes could again happen directly, which allowed for efficient selection. However, the performance of heterogeneous teams that evolved with individual selection remained lower than that of homogeneous teams that evolved with team selection (P < 0.015 and P < 0.003, respectively, Fig. 11). This may have been because heterogeneous teams had to solve a more complex optimization task.

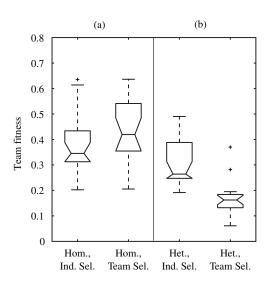


Fig. 11. Task 3—Altruistic cooperative foraging without disparities in genome evaluation and without credit assignment problems. (a) Homogeneous teams evolved with one evaluation per team (instead of 10) and (b) heterogeneous teams evolved with 100 agents per population (instead of 1000). The performance of heterogeneous teams evolved with individual selection was higher than the performance of heterogeneous teams evolved with evolved with team selection, but did not reach that of homogeneous teams. For boxplot explanations, see Fig. 5.

In this second additional set of experiments, the performance of homogeneous teams that evolved with individual selection and homogeneous teams that evolved with team selection did not differ (P = 0.133).

Heterogeneous teams evolved with team selection performed worse than all other evolutionary conditions due to inefficient selection (all three P < 0.001).

Importantly, the altruistic cooperative foraging task led to the evolution of a different foraging strategy in heterogeneous teams evolved with individual selection than in the other three evolutionary conditions (Fig. 10). A possible reason is that cooperation to collect large tokens now implied a cost for individuals. To test this hypothesis, we performed additional experiments with this evolutionary condition. First we repeated the experiments with a setup identical to that of task 3, i.e.,

<sup>&</sup>lt;sup>4</sup>An electronic supplement for this paper is available online at http://lis.epfl.ch/documentation.php

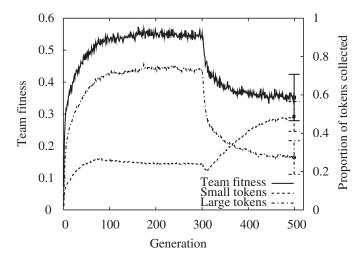


Fig. 12. Task 3—Altruistic Cooperative Foraging in heterogeneous teams evolved with individual selection. For the first 300 generations, individual contributions to the cooperative foraging of large tokens were known (no credit assignment problems). From generation 300 onward, individual contributions to the foraging of large tokens were presumed unknown (credit assignment problems). The introduction of credit assignment problems led to the rapid collapse of cooperation and a decrease in team fitness.

with 1000 agents per population and 10 evaluations per team, but with known individual contributions to large token foraging, i.e., a situation without credit assignment problems. Each of the four large tokens awarded five fitness points to each of the two transporting robots, rather than one fitness point to each of the ten team members. Then, at generation 300, we changed the fitness assignment and assumed unknown individual contributions to large token foraging, i.e., a situation with credit assignment problems. Each of the four large tokens awarded 1 fitness point to each team member, irrespective of their participation in token foraging.

This change in fitness assignment resulted in a drastic change in foraging strategy (Fig. 12). While at generation 300 heterogeneous teams that evolved with individual selection collected a significantly higher proportion of large tokens than small tokens (P < 0.001), at generation 500 they collected a significantly lower proportion of large than small tokens (P <0.001). As a direct result of this change, team performance decreased significantly (P < 0.001) between generation 300 and generation 500. This was because after the introduction of credit assignment problems, fitness points gained from large tokens were assigned to all team members, and therefore individuals collecting small tokens gained a fitness advantage over their team mates. This led to the selection of individuals that foraged for small tokens and resulted in fewer and fewer individuals foraging for large tokens. The observed drop in team fitness also implies a drop in average individual fitness. This illustrates that fitness is a relative measure of performance and therefore evolution selects for performance increase relative to the performance of competitors, rather than for absolute performance. The simplicity of the neural network controllers did not allow individuals to accurately discriminate large and small tokens, which explains the incomplete collapse of large token foraging.

In contrast, the foraging strategy in homogeneous teams that evolved with individual selection and in homogeneous teams that evolved with team selection was not affected by the costs implied in large token foraging (see Figs. S5 and S6 in the supplementary material<sup>5</sup>). This was because relative fitness differences between team members could not have an influence on the selection of genomes when individuals were genetically identical.

Foraging strategy in heterogeneous teams that evolved with team selection was not affected by the costs implied in large token foraging (see Figs. S5 and S6 in the supplementary material<sup>5</sup>). This was because relative fitness differences between team members did not have an influence on selection of genomes when selection acted at the level of the team.

#### V. CONCLUSION

This paper provides an experimental demonstration of how the choice of genetic team composition and level of selection influences the performance of MAS in tasks with varying levels of cooperation that do not provide a benefit for specialization. We have identified three different types of multiagent tasks depending on the amount of cooperation required between team members. Our results demonstrate that different combinations of genetic team composition and level of selection lead to significant performance differences. No combination achieved optimal performance in all three types of task.

We have identified and studied three different types of multiagent tasks depending on the amount of cooperation required between team members.

In tasks that did not require cooperation, heterogeneous teams that evolved with individual level selection achieved the highest team performance. Team heterogeneity allowed evaluation of a high number of different genomes in parallel, and individual selection allowed efficient selection of good genomes. However, these teams performed poorly in tasks that required cooperation and in tasks with credit assignment problems.

For multiagent tasks that required cooperation, the highest team performance was achieved by homogeneous teams. These teams led to efficient cooperation between team members and they were not affected by credit assignment problems or costs associated with cooperation. Our results suggest that homogeneous teams are a safe choice in tasks that do not benefit from specialization when the requirements for agent cooperation are difficult to estimate. Compared to heterogeneous teams, homogeneous teams evaluate less genomes, which may result in premature convergence to suboptimal solutions. Our experimental results indicate that a simple way to prevent this problem is to use populations made of a large number of homogeneous teams.

Heterogeneous teams evolved with team selection were inefficient at selecting for good performance in all three types of tasks studied here and therefore cannot be recommended for cooperative tasks that do not require specialization.

Converging evidence towards these guidelines was recently found in a study on the evolutionary conditions for the emergence of communication in societies of robots with different

<sup>5</sup>An electronic supplement for this paper is available online at http://lis.epfl.ch/documentation.php

morphologies and sensing abilities than those described in this paper [17].

However, it should be noted that evidence from both studies has two notable limitations. First, they did not address tasks that benefit from specialization in addition to cooperation. There is evidence that behavioral heterogeneity can lead to significant performance advantages for such tasks [29], [63], [85]–[88]. Second, they did not consider teams with intermediate genetic similarity. Biological research has shown that such teams can overcome individual fitness costs of cooperation [55], [89], thus combining the best of both worlds—enhanced genetic diversity with readiness to adopt altruistic behaviors. A good understanding of those conditions will require significant further research.

#### REFERENCES

- G. Baldassarre, S. Nolfi, and D. Parisi, "Evolving mobile robots able to display collective behavior," *Artificial Life*, vol. 9, no. 3, pp. 255–267, 2003.
- [2] S. Nolfi and D. Floreano, *Evolutionary Robotics*. Bradford, U.K.: MIT Press, 2000.
- [3] A. Agogino, K. Stanley, and R. Miikkulainen, "Online interactive neuroevolution," *Neural Process. Lett.*, vol. 11, no. 1, pp. 29–38, 2000.
- [4] D. Andre, "The automatic programming of agents that learn mental models and create simple plans of action," in *Proc. 14th Int. Joint Conf. Artificial Intell.* '95, vol. 1. pp. 741–747.
- [5] D. Andre and A. Teller, *RoboCup-98: Robot Soccer World Cup II* (Evolving Team Darwin United). vol. 1604, Berlin/Heidelberg, Germany: Springer-Verlag, pp. 346–351, 1999.
- [6] G. Baldassarre, S. Nolfi, and D. Parisi, "Evolution of collective behavior in a team of physically linked robots," in *Applications in Evolutionary Computing*, Berlin, Germany: Springer-Verlag, 2003, pp. 581–592.
- [7] G. Baldassarre, D. Parisi, and S. Nolfi, "Coordination and behavior integration in cooperating simulated robots," in *Proc. 8th Int. Conf. Simulation Adaptive Behavior From Animals to Animats*, Cambridge, MA: MIT Press, 2003, pp. 385–394.
- [8] C. Baray, "Evolving cooperation via communication in homogeneous multiagent systems," in *Proc. Intell. Inform. Syst.*, 1997, Grand Bahama Island, Bahamas, Dec. 1997, pp. 204–208.
- [9] C. Baray, "Effects of population size upon emergent group behavior," *Complexity Int.*, vol. 6, 1998.
- [10] J. K. Bassett and K. A. De Jong, "Evolving behaviors for cooperating agents," in *Proc. 12th Int. Symp. Found. Intell. Syst. (ISMIS 2000)*, Berlin/Heidelberg, Germany: Springer-Verlag, pp. 157–165.
- [11] J. C. Bongard, "The legion system: A novel approach to evolving hetrogeneity for collective problem solving," *Lecture Notes Comput. Sci.*, LNCS vol. 1802/2004, pp. 16–28, 2000.
- [12] H. Botee and E. Bonabeau, "Evolving ant colony optimization," Advances Complex Syst., vol. 1, no. 2/3, pp. 149–159, 1998.
- [13] B. Bryant and R. Miikkulainen, "Neuroevolution for adaptive teams," in Proc. 2003 Congr. Evol. Comput., vol. 3. Dec. 2003, pp. 2194–2201.
- [14] L. Bull and O. Holland, "Evolutionary computing in multiagent environments: Eusociality," in *Proc. Annu. Conf. Genetic Programming*, 1997, pp. 347–352.
- [15] R. J. Collins and D. R. Jefferson, "Antfarm: Towards simulated evolution," in *Artificial Life II*, Redwood City, CA: Addison-Wesley, 1992, pp. 579–601.
- [16] M. Dorigo, V. Trianni, E. Scedilahin, R. Gross, T. H. Labella, G. Baldassarre, S. Nolfi, J. L. Deneubourg, F. Mondada, D. Floreano, and L. M. Gambardella, "Evolving self-organizing behaviors for a swarm-bot," *Autonomous Robots*, vol. 17, no. 2-3, pp. 223–245, 2004.
- [17] D. Floreano, S. Mitri, S. Magnenat, and L. Keller, "Evolutionary conditions for the emergence of communication in robots," *Current Biology*, vol. 17, no. 6, pp. 514–519, 2007.
- [18] R. Gross and M. Dorigo, "Cooperative transport of objects of different shapes and sizes," *Lecture Notes Comput. Sci.*, vol. 3172, pp. 106–117, 2004.
- [19] R. Gross and M. Dorigo, "Evolving a cooperative transport behavior for two simple robots," *Lecture Notes Comput. Sci.*, vol. 2936, pp. 305–316, 2004.

- [20] A. Hara and T. Nagao, "Emergence of cooperative behavior using ADG; Automatically defined groups," in *Proc. 1999 Genetic Evol. Comput. Conf.*, San Mateo, CA: Morgan Kaufmann, pp. 1038–1046.
- [21] T. Haynes, R. Wainwright, S. Sen, and D. Schoenefeld, "Strongly typed genetic programming in evolving cooperation strategies," in *Proc. 6th Int. Conf. Genetic Algorithms*, San Mateo, CA: Morgan Kaufman, 1995, pp. 271–278.
- [22] T. Haynes and S. Sen, "Crossover operators for evolving a team," in Proc. 2nd Annu. Conf. Genetic Programming 1997, San Mateo, CA: Morgan Kaufmann, Stanford, CA: Stanford University, pp. 162–167.
- [23] J. Lohn and J. Reggia, "Discovery of self-replicating structures using a genetic algorithm," in *Proc. IEEE Int. Conf. Evol. Comput. 1995*, vol. 2. Dec. 1995, pp. 678–683.
- [24] S. Luke and L. Spector, "Evolving teamwork and coordination with genetic programming," in *Proc. Genetic Programming 1996*, Stanford, CA: Stanford Univ. Press, pp. 150–156.
- [25] S. Luke, C. Hohn, J. Farris, G. Jackson, and J. Hendler, "Co-evolving soccer softbot team coordination with genetic programming," in *Proc. 1st Int. Workshop RoboCup, IJCAI-97*, Nagoya, Aichi: Springer-Verlag, pp. 214–222.
- [26] S. Luke, "Genetic programming produced competitive soccer softbot teams for robocup 97," in *Proc. 3rd Annu. Conf. Genetic Programming* 1998, San Mateo, CA: Morgan Kaufmann, pp. 214–222.
- [27] D. Marocco and S. Nolfi, "Self-organization of communication in evolving robots," in *Proc. 10th Int. Conf. Artificial Life: Alife*, Boomington, MN: MIT Press, 2006, pp. 199–205.
- [28] T. Miconi, "When evolving populations is better than coevolving individuals: The blind mice problem," in *Proc. 18th Int. Joint Conf. Artificial Intell.*, San Mateo, CA: Morgan Kaufmann, 2003, pp. 647–652.
- [29] M. Quinn, "A comparison of approaches to the evolution of homogeneous multirobot teams," in *Proc. 2001 Congr. Evol. Comput.*, Seoul, Korea: IEEE, pp. 128–135.
- [30] M. Quinn, L. Smith, G. Mayley, and P. Husbands, "Evolving controllers for a homogeneous system of physical robots: Structured cooperation with minimal sensors," *Philosophical Trans. Royal Soc. London Ser. A: Math. and Physical and Eng. Sci.*, vol. 361, pp. 2321–2344, 2003.
- [31] S. Raik and B. Durnota, "The evolution of sporting strategies," in *Complex System: Mechanisms of Adaption*, Amsterdam, Netherlands: IOS Press, 1994, pp. 85–92.
- [32] M. Richards, D. Whitley, J. Beveridge, T. Mytkowicz, D. Nguyen, and D. Rome, "Evolving cooperative strategies for UAV teams," in *Proc.* 2005 Conf. Genetic Evol. Comput., pp. 1721–1728.
- [33] A. Robinson and L. Spector, "Using genetic programming with multiple data types and automatic modularization to evolve decentralized and coordinated navigation in multiagent systems," in *Proc. Int. Soc. Genetic Evol. Comput., Late-Breaking Papers Genetic Evol. Comput. Conf.* '02.
- [34] G. Saunders and J. Pollack, "The evolution of communication schemes over continuous channels," in *Proc. 4th Int. Conf. Adaptive Behavior From Animals Animats*, 1996.
- [35] V. Trianni, R. Gross, T. H. Labella, E. Sahin, and M. Dorigo, "Evolving aggregation behaviors in a swarm of robots," *Lecture Notes Comput. Sci.*, vol. 2801, pp. 865–874, 2003.
- [36] V. Trianni, S. Nolfi, and M. Dorigo, "Cooperative hole-avoidance in a swarm-bot," *Robotics Autonomous Syst.*, vol. 54, pp. 97–103, 2006.
- [37] A. S. Wu, A. C. Schultz, and A. Agah, "Evolving control for distributed micro air vehicles," in *Proc. IEEE Conf. Comput. Intell. Robotics*, Monterey, CA, 1999, pp. 174–179.
- [38] N. Zaera, C. Cliff, and J. Bruten, "(Not) evolving collective behaviors in synthetic fish," in *Proc. 4th Int. Conf. Simulation Adaptive Behavior* '96, vol. 4. pp. 635–644.
- [39] M. Mirolli and D. Parisi, "How can we explain the emergence of a language that benefits the hearer but not the speaker," *Connection Sci.*, vol. 17, no. 3, pp. 307–324, 2005.
- [40] A. Martinoli, "Swarm intelligence autonomous collective robotics: From tools to analysis and synthesis distributed control strategies," Ph.D. dissertation, Ecole Polytechnique Fédérale de Lausanne (EPFL), Lausanne, Switzerland, 1999.
- [41] A. Agah and G. A. Bekey, "Phylogenetic and ontogenetic learning in a colony of interacting robots," *Autonomous Robots*, vol. 4, no. 1, pp. 85– 100, 1997.
- [42] A. Agah and K. Tanie, "Robots playing to win: evolutionary soccer strategies," in *Proc. IEEE Int. Conf. Robotics Automation* '97, vol. 1. Albuquerque, NM, Apr. 1997, pp. 632–637.
- [43] R. Bianco and S. Nolfi, "Toward open-ended evolutionary robotics: Evolving elementary robotic units able to self-assemble and selfreproduce," *Connection Sci.*, vol. 16, no. 4, pp. 227–248, 2004.

- [44] A. Cangelosi and D. Parisi, "The emergence of a 'language' in an evolving population of neural networks," *Connection Sci.*, vol. 10, no. 2, pp. 83–97, 1998.
- [45] C. Reynolds, "An evolved, vision-based behavioral model of coordinated group motion," in *From Animals to Animats*. Cambridge, MA: MIT Press, 1993, pp. 384–392.
- [46] G. M. Werner and M. G. Dyer, "Evolution of herding behavior in artificial animals," in *Proc. 2nd Int. Conf. From Animals to Animats: Simulation Adaptive Behavior*. Cambridge, MA: MIT Press, 1992, pp. 393–399.
- [47] G. M. Werner and M. G. Dyer, "Evolution of communication in artificial organisms," in *Artificial Life II*, Redwood City, CA: Addison-Wesley, 1992, pp. 659–687.
- [48] C. R. Ward, F. Gobet, and G. Kendall, "Evolving collective behavior in an artificial ecology," *Artificial Life*, vol. 7, no. 2, pp. 191–209, 2001.
- [49] K. Stanley, B. Bryant, and R. Miikkulainen, "Real-time neuroevolution in the NERO video game," *IEEE Trans. Evol. Comput.*, vol. 9, no. 6, pp. 653–668, Dec. 2005.
- [50] S. G. Ficici, R. A. Watson, and J. B. Pollack, "Embodied evolution: A response to challenges in evolutionary robotics," in 8th Eur. Workshop Learning, 1999, pp. 14–22.
- [51] E. D. V. Simoes and D. A. C. Barone, "Predation: An approach to improving the evolution of real robots with a distributed evolutionary controller," in *Proc. 2002 IEEE Int. Conf. Robotics Automation*, vol. 1. May 2002, pp. 664–669.
- [52] L. Spector, J. Klein, C. Perry, and M. Feinstein, "Emergence of collective behavior in evolving populations of flying agents," *Lecture Notes Comput. Sci.*, vol. 2723, pp. 61–73, 2003.
- [53] L. Spector, J. Klein, C. Perry, and M. Feinstein, "Emergence of collective behavior in evolving populations of flying agents," *Genetic Programming and Evolvable Mach.*, vol. 6, no. 1, pp. 111–125, 2005.
- [54] R. A. Watson, S. G. Ficici, and J. B. Pollack, "Embodied evolution: Distributing an evolutionary algorithm in a population of robots," *Robotics Autonomous Syst.*, vol. 39, pp. 1–18, 2002.
- [55] A. S. Griffin, S. A. West, and A. Buckling, "Cooperation and competition in pathogenic bacteria," *Nature*, vol. 430, no. 7003, pp. 1024–1027, 2004.
- [56] C. Hauert, S. De Monte, J. Hofbauer, and K. Sigmund, "Replicator dynamics for optional public good games," *J. Theoretical Biology*, vol. 218, no. 2, pp. 187–194, 2002.
- [57] M. A. Potter and D. V. De Jong, "A cooperative coevolutionary approach to function optimization," *Lecture Notes Comput. Sci.*, vol. 866, pp. 249–257, 1994.
- [58] H. J. Blumenthal and G. B. Parker, "Co-evolving team capture strategies for dissimilar robots," in *Proc. AAAI 2004 Symp. Artificial Multiagent Learning*, Washington, D.C.
- [59] M. A. Potter and D. V. De Jong, "Cooperative coevolution: An architecture for evolving coadapted subcomponents," *Evol. Comput.*, vol. 8, no. 1, pp. 1–29, 2000.
- [60] R. P. Wiegand, W. C. Liles, and D. V. DeJong, "An empirical analysis of collaboration methods in cooperative coevolutionary algorithms," in *Proc. Genetic Evol. Comput. Conf.*, 2001, pp. 1235–1242.
- [61] K.-C. Jim and C. L. Giles, "Talking helps: Evolving communicating agents for the predator-prey pursuit problem," *Artificial Life*, vol. 6, no. 3, pp. 237–254, 2001.
- [62] M. Quinn, L. Smith, G. Mayley, and P. Husbands, "Evolving teamwork and role-allocation with real robots," in *Proc. 8th Int. Conf Artificial Life*, Cambridge, MA: MIT Press, 2002, pp. 302–311.
- [63] J. C. Bongard, "Reducing collective behavioral complexity through heterogeneity," in *Proc. 7th Int. Conf. Artificial Life*, 2000, pp. 327–336.
- [64] X. Yao, "Evolving artificial neural networks," *Proc. IEEE*, vol. 87, no. 9, pp. 1423–1447, Sep. 1999.
- [65] D. C. Queller, "Genetic relatedness in viscous populations," *Evol. Ecology*, vol. 8, no. 1, pp. 70–73, 1994.
- [66] S. A. West, M. G. Murray, C. A. Machado, A. S. Griffin, and E. A. Herre, "Testing Hamilton's rule with competition between relatives," *Nature*, vol. 409, pp. 510–513, 2001.
- [67] W. D. Hamilton, "The Genetical Evolution of Social Behavior I+II," J. Theoretical Biology, vol. 7, pp. 1–52, 1964.

- [68] L. Lehmann and L. Keller, "The evolution of cooperation and altruism: A general framework and a classification of models." *J. Evol. Biology*, vol. 19, no. 5, pp. 1365–1376, 2006.
- [69] A. Agogino and K. Tumer, "Efficient evaluation functions for multi-rover systems," *Lecture Notes Comput. Sci.*, vol. 3102, pp. 1–11, 2004.
- [70] J. Grefenstette, "Credit assignment in rule discovery systems based on genetic algorithms," *Mach. Learning*, vol. 3, no. 2, pp. 225–245, 1988.
  [71] M. Minsky, "Steps toward artificial intelligence," in *Proc. Inst. Radio*
- [71] M. Minsky, "Steps toward artificial intelligence," in *Proc. Inst. Radio Eng.*, vol. 49. New York, NY: McGraw-Hill, 1961, pp. 8–30.
- [72] L. Panait and S. Luke, "Cooperative multiagent learning: The state of the art," Autonomous Agents and Multiagent Syst., vol. 11, no. 3, pp. 387– 434, 2005.
- [73] L. Keller, *Levels of Selection in Evolution*. Princeton, NJ: Princeton University Press, 1999.
- [74] S. West, A. Gardner, and A. Griffin, "Altruism," *Current Biology*, vol. 16, no. 13, pp. 482–483, 2006.
- [75] W. D. Hamilton, "Altruism and related phenomena, mainly in social insects," Annu. Rev. Ecology and Syst., vol. 3, no. 1, pp. 193–232, 1972.
- [76] T. Balch, "The impact of diversity on performance in multirobot foraging," in *Proc. Int. Conf. Autonomous Agents*, Seattle, WA, 1999, pp. 92–99.
- [77] Y. U. Cao, A. S. Fukunage, and A. B. Kahng, "Cooperative mobile robotics: Antecedents and directions," in *Proc. Expanded Version 1995 IEEE/RSJ IROS Conf.*, 1997, pp. 7–27.
- [78] M. J. B. Krieger, J. B. Billeter, and L. Keller, "Ant-like task allocation and recruitment in cooperative robots," *Nature*, vol. 406, no. 6799, pp. 992–995, 2000.
- [79] J. L. Reyes Lopez, "Optimal foraging in seed-harvester ants: Computeraided simulation," *Ecology*, vol. 68, no. 6, pp. 1630–1633, 1987.
- [80] J. F. A. Traniello, "Foraging strategies of ants," Annu. Rev. Entomology, vol. 34, no. 1, pp. 191–210, 1989.
- [81] G. Caprari, "Autonomous micro-robots: Applications and limitations," Ph.D. dissertation, Ecole Polytechnique Fédérale de Lausanne (EPFL), Lausanne, Switzerland, 2003.
- [82] S. Magnenat and M. Waibel. (2007). Enki, a Fast Physics-Based 2-D Simulator [Online]. Available: http://teem.epfl.ch
- [83] S. Magnenat, M. Waibel, and A. Beyeler. (2007). Teem, an Open Evolutionary Framework [Online]. Available: http://teem.epfl
- [84] R. McGill, J. Tukey, and W. Larsen, "Variations of box plots," *Amer. Statist.*, vol. 32, no. 1, pp. 12–16, 1978.
- [85] T. Balch, "Behavioral diversity in learning robot teams," Ph.D. dissertation, Georgia Institute of Technology, Atlanta, GA, 1998.
- [86] M. Waibel, D. Floreano, S. Magnenat, and L. Keller, "Division of labour and colony efficiency in social insects: Effects of interactions between genetic architecture, colony kin structure and rate of perturbations." in *Proc. Royal Soc. B*, vol. 273. 2006, pp. 1815–1823.
- [87] M. A. Potter, L. A. Meeden, and A. C. Schultz, "Heterogeneity in the coevolved behaviors of mobile robots: The emergence of specialists," in *Proc. 7th Int. Conf. Artificial Intell.*, 2001, pp. 1337–1343.
- [88] D. Tarapore, D. Floreano, and L. Keller, "Influence of the level of polyandry and genetic architecture on division of labour," in *Proc. 10th Int. Conf. Simulation Synthesis Living Syst.*, 2006, pp. 358–364.
- [89] A. S. Griffin and S. A. West, "Kin selection: Fact and fiction," *Trends Ecol. Evol.*, vol. 17, no. 1, pp. 15–21, 2002.



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