ORIGINAL ARTICLE

Efects of excessive elitism on the evolution of artifcial creatures with NEAT

Siti Aisyah Binti Jaafar1 · Reiji Suzuki¹ · Satoru Komori1 · Takaya Arita¹

Received: 5 July 2023 / Accepted: 6 March 2024 / Published online: 3 April 2024 © International Society of Artifcial Life and Robotics (ISAROB) 2024

Abstract

This paper proposes a simple method based on a novel use of elitism to increase the population size of artifcial creatures while minimizing evaluation cost. This can contribute to preventing premature convergence of the population. We propose the "Excessive Elitism (EE)" method by modifying elitism in HyperNEAT (Hypercube-based NeuroEvolution of Augmenting Topologies), which is an evolutionary algorithm commonly used to evolve genotype [i.e., Compositional Pattern Producing Network (CPPN)] of artifcial creatures. In EE, the evaluated ftness of best-ft individuals will be succeeded and reused instead of being re-evaluated during subsequent ftness evaluations, thereby reducing the evaluation cost if the elite size is excessive. Notably, EE also disables speciation and ftness sharing, serving to simplify the population structure and reduce complexity. In a 3D multi-agent environment, we evolved the morphology and behavior of artifcial creatures with a simple target approach task. We assumed a baseline case $(EE (2, 20))$ in which a small population size was used due to the strong limitation of the evaluation cost and adopted a normal small elite size. This often led to premature convergence of the population to suboptimal individuals who could not reach the target. However, with the application of EE, the population was capable of evolving to reach the target, maintaining an evaluation cost comparable to EE (2, 20). We demonstrate that EE method serves as a simpler alternative to speciation for diversity preservation, capable of enhancing both the average and optimal ftness of a population, thus preventing premature convergence at a minimal evaluation cost. Further research in complex environments is required to fully uncover the potential and limitations of this method.

Keywords Artifcial creatures · Artifcial life · Elitism · Evaluation cost · 3D multi-agent simulation

1 Introduction

An approach to evolving artificial creatures has gained significant interest in scientific (e.g., evo-devo $[1, 2]$ $[1, 2]$ $[1, 2]$ $[1, 2]$) and engineering research (e.g., design of soft robots $[1-3]$ $[1-3]$). In particular, this approach has proven useful for understanding the interactions between ecological, developmental, and evolutionary processes known as eco-evo-devo. Furthermore, it has provided clarity on the principles that govern

 \boxtimes Siti Aisyah Binti Jaafar binti.jaafar.siti.aisyah.v4@s.mail.nagoya-u.ac.jp the relationship between environmental complexity and the evolution of morphological and behavioral traits. We aim to discuss the evolution of development and niche construction within a 3D multi-agent environment. To achieve this, we will utilize the framework by Chiba et al. [\[3](#page-10-2)] which enables the evolution of creatures' behaviors toward environmental modifcation.

However, incorporating various factors related to the above topics into the model has signifcantly increased the evaluation cost for the physical simulation. As a result, evolutionary experimentation grown in complexity and computational expense. This is largely due to the substantial cost associated with ftness evaluation, which relies on a physical simulation of complex creatures' behaviors, complexity that emerges from the interactions among the creatures. This has been a signifcant challenge in the evolution of artifcial creatures, thoroughly explored in several previous studies [[4,](#page-10-3) [5](#page-10-4)]. Consequently, a small population size is often used as an

This work was presented in part at the joint symposium of the 28th International Symposium on Artifcial Life and Robotics, the 8th International Symposium on BioComplexity, and the 6th International Symposium on Swarm Behavior and Bio-Inspired Robotics (Beppu, Oita and Online, January 25–27, 2023).

¹ Graduate School of Informatics, Nagoya University, Nagoya, Japan

ad hoc way to resolve the problem, although it may lead to the premature convergence of the population to local optima.

To overcome the problem, a number of studies have expressed confdence that the implementation of elitism can lead to performance improvements. For example, in the feld of soft robot evolution, La Cava and Moore proposed search driver algorithms aimed at fnding optimal solutions [\[6](#page-10-5)]. Elitism, a methodology that involves either preserving the individual with the best ftness for the next generation or replacing the least ft individual with a previous high-ftness individual, was implemented in some of these algorithms. The study's comparative analysis of the algorithms demonstrated that simulations introducing some form of elitism signifcantly contributed to the discovery of locomoting soft robot morphologies.

Methenitis et al. [[7\]](#page-10-6) also suggested that implementing the ftness-elitism technique using NeuroEvolution of Augmenting Topologies (NEAT) [\[8](#page-10-7)], which is a popular evolutionary algorithm for evolving complex artifcial neural networks, improved the performance in evolving soft robots. The study began by experimenting with two exclusive methods. First, it involved a novelty-search method, which promotes diversity by rewarding novel individuals regardless of their ftness. Second, it implemented ftness elitism, where the best individuals are selected based on their ftness, thereby ensuring the most optimal genetic material is passed from one generation to the next. The study later revealed that by integrating both methods, the ft and unique individual was carried over to subsequent generations, potentially resulting in the production of ofspring that are both novel and ft. Their fndings emphasized that the combination of methodologies played a critical part in advancing the evolution of soft robots. Moreover, these fndings suggests that the presence of elitism signifcantly enhances the evolution performance of artifcial creatures. Beyond simply improving ftness through elitism, it is also crucial to consider the potential benefts it provides in reducing evaluation costs linked to complex components inherent in evolutionary simulations of creatures.

Elitism, as mentioned, has exhibited signifcant potential in enhancing the performance of artifcial creatures' evolution. In coherence with this, an interesting aspect to consider is the role speciation plays in evolution when using NEAT. Speciation based on the affinity of phenotypes and ftness sharing among conspecifc individuals serve as key mechanisms for NEAT to maintain the diversity of individuals within a small population. However, the implementation of speciation requires a metric to measure the phenotypic distance between individuals. Consequently, the population structure may become complicated, subsequently making the population dynamics more complex compared to the case with a single species. While the efective combination of methods and maintaining diversity in species presents complexities, it makes the evolutionary process an intriguing feld to explore further.

Given these complexities, this paper introduces a novel approach called "Excessive Elitism (EE)" derived from the evolutionary algorithm Hypercube-based NeuroEvolution of Augmenting Topologies (HyperNEAT) [[9\]](#page-10-8), frequently used to evolve the genotype (i.e., Compositional Pattern Producing Network (CPPN) [[10](#page-10-9)]) of artificial creatures. The principal mechanism of EE is that the evaluated ftness of best-ft individuals will be succeeded and reused instead of being re-evaluated during subsequent ftness evaluation. This contributes to a reduced evaluation cost within a large population, particularly when applying an excessively large elite size. Consequently, the EE method presents a straightforward alternative to speciation and ftness sharing to prevent premature convergence [\[11](#page-10-10), [12](#page-10-11)] of the population. This enables an increase in the population size while maintaining relatively low evaluation costs.

In this study, our focus is on evolution of the genotypes of a rigid-bodied creatures in a 3D-multi-agent environment, a situation typically resulted in the occurrence of premature convergence to local optima. We conducted experiments using varying population sizes whilst maintaining a consistent number of evaluated individuals. The outcome of our experiments revealed that employing EE in a large, single population proved more efective than in smaller populations, even when maintaining an equivalent number of evaluated individuals for each generation. Moreover, the fndings pointed towards a potential optimal confguration for population size. Notably, EE showcased a morphological diversity comparable to standard NEAT-evolved populations that involved speciation. This suggests that EE is capable of evolving a population with an adaptability level similar to those evolved by NEAT, while concurrently preserving a less complex population structure.

2 Model

We use a framework for evolving artificial creatures where the morphology of individuals develops within a 3D-multiagent environment. This is based on the Python modulebased physics engine, PyBullet, employed to analyze eco-evo-devo in evolving artifcial creatures (Fig. [1](#page-2-0)). The HyperNEAT (Hypercube-based NeuroEvolution of Augmenting Topologies) is adopted as the evolutionary algorithm to evolve the genotype (i.e., the Compositional Pattern Producing Network or CPPN) of artifcial creatures. Hyper-NEAT is an evolutionary algorithm used for evolving complex neural networks, represented by the CPPN in the model [[10\]](#page-10-9). We utilize this algorithm to evolve the CPPN as the genotype of each artifcial creature, used in their morphological development and behavioral generation during the

Fig. 1 The lifetime of a creature illustrated through a repeating process that involves two key steps: (1) the generation of behavior through connection weight settings (**a**), and (2) morphological development that takes place via new block additions (**b**), with four occurrence events

evolutionary process. We assume that a single large CPPN determined by the genotype of a creature represents both CPPNs as its sub-networks. Accordingly, the genotype of a creature generates two neural networks: one for determining the morphological development and another for determining the behavioral generation process. These components are depicted in Fig. [3](#page-2-1) and will be discussed in greater detail in the sections that follow.

2.1 Morphological development

Figure [1](#page-2-0) illustrates the lifetime of a creature within a population. Each rigid-bodied creature is comprised of rectangular blocks interconnected via hinges (Fig. [2](#page-2-2)(ii)–(iii)). The initial body, composed of a single block, is positioned at the origin of the substrate space—the space where developmental events occur (Fig. $2(i)$ $2(i)$ –(iii)). The creature's morphology develops through the addition of new blocks to pre-existing body blocks, in several fxed time steps throughout its lifetime.

Fig. 3 The neural network architecture of the genotype of an artifcial creature, as represented by the Compositional Pattern Producing Network (CPPN). (Left) the CPPN is utilized for morphological develop-

ment via the addition of a new block. (Right) the CPPN is employed to calculate connection weights for nodes in the three-layer behavioral generation network in substrate space

Each developmental event involves the addition of a new block to each candidate position (yellow circle) on the existing body blocks. Every block has three candidate positions. The genotype, CPPN (Fig. [3b](#page-2-1) left), receives three inputs: the elapsed time from the birth, the central coordinate of the focal block on the substrate space, and the coordinate of a candidate position of the focal block. This generates four outputs: determining if a new block is added or not, the length of the long sides of the new block, hinge direction, and deciding between a fxed hinge and fexible joint. The size of each new block incrementally increases from 0 to a specifed length, up until the end of the developmental event. This process is spread over a total of 500-time steps.

An example illustrated in Fig. [2\(](#page-2-2)i)–(iii) demonstrates how the process begins with a single block serving as the initial morphology (i). Subsequently, a new block grows at a right-candidate position (ii), creating candidate positions for additional blocks to grow during subsequent developmental events (iii).

2.2 Behavioral generation

After the completion of each morphological developmental event, the behavioral generation network (Fig. [3](#page-2-1) (right)) is constructed by the genotype or updated in accordance with the creature's morphology changes. This network structure comprises three layers: input, intermediate, and output. Each block has a centrally located radar sensor (Fig. [4\)](#page-3-0) shown in purple, and a corresponding radar node that receive the sen-sory information from the substrate space (Fig. [3](#page-2-1) (middle)). The radar sensor calculates the distance from the creature's focal block to the target, utilizing the block's angle and orientation relative to the target object in the evaluation feld as depicted in Fig. [4](#page-3-0). Consequently, the radar node receives the distance to the target in that specifc direction and provides the

Fig. 4 Illustration of the radar sensor method. Here, we consider the longer side of the rectangular block as the 'front side'. The angle of rotation derived from line (a) illustrates the current orientation of the block

input to the behavioral generation network, allowing the neural network to attempt to minimize this distance by manipulating the fexible hinges of the creature. Each hinge between blocks has a corresponding hinge node (red), and there are several intermediate nodes scattered around the creature on the substrate space (Fig. [2](#page-2-2)iii).

We assume that the radar, intermediate, and hinge nodes are located on the input, intermediate, and output layers of the behavioral generation network, respectively. For instance, in the network depicted in Fig. [3](#page-2-1) (middle), there are 2 radar sensor nodes (input layer), 7 intermediate nodes (intermediate layer), and 1 hinge node (output layer). Connection exists among the nodes between the input-intermediate, intermediate-output, and input–output layers. Each linkage possesses its own connection weight, which is determined by the CPPN receiving the coordinates of the source and destination nodes pair on the substrate space. Each node within the output layer conveys the angle of the corresponding hinge, adjusting the actual angle of the physical hinge to match this output value. Consequently, the behavioral generation network can produce diverse behavioral patterns in the creature, according to the sensory input received from the radars.

2.3 Task

Figure [5](#page-3-1) illustrates a feld for ftness evaluation. We conducted experiments involving a locomotion task to examine the implications of excessive elitism on evolution. The artifcial creatures were positioned in a circular manner, surrounding a central target (cube) placed on a fat surface. Fitness was defned as the distance traveled by creatures from the initial position towards the target.

Specifcally, each creature's ftness is calculated using the equation provided in Eq. ([1\)](#page-3-2)

$$
Fitness = \begin{cases} d_i - d_{io}, (d_{io} \le d_i) \\ 0, (d_{io} > d_i) \end{cases}.
$$
 (1)

In this equation, d_i denotes the initial distance between the creature and the target, a value constant in all experiments,

Fig. 5 Fitness evaluation feld. **A** presents a top-down perspective, featuring the initial placement of creatures, who are arranged in a circular pattern, with an initial distance of 60 separating them from the target. **B** provides a lateral view of the evaluation feld

which is 60. d_{io} is the current distance between the creature and the target object. From Eq. ([1\)](#page-3-2), we assume that if the distance between the creature and the target reaches or falls below 60, indicating the creature is either stationary or approaching the target, the ftness corresponds to the distance covered. However, if the creature moves away from its initial position, resulting in a negative total distance, the ftness is considered zero.

2.4 Evolution by HyperNEAT

In this section, we delve into how the evolution of the artificial creatures in this study was conducted using Hyper-NEAT (Hypercube-based NeuroEvolution of Augmenting Topologies) [[9\]](#page-10-8), an evolutionary algorithm frequently used to evolve the genotype of artifcial creatures, namely the Compositional Pattern Producing Network (CPPN) [[14](#page-11-0)]. HyperNEAT evolves the genome, a connective CPPN that generates a substrate by computing connection weights to solve a problem.

The selection method employed in HyperNEAT is derived from its underlying evolutionary algorithm, NEAT. According to NEAT, a solution to a problem evolves through the ftness function, which computes a singular number indicative of the quality of each individual genome (60); a higher score signifes enhanced ability. The algorithm proceeds through a pre-determined number of generations (400), each of which is generated through the reproduction and mutation of the fttest individuals from the preceding generation. The processes of reproduction and mutation may introduce additional nodes and/or connections to the genomes.

The variation operators in HyperNEAT play a crucial role in introducing genetic diversity into the population of genomes, which represents ANNs, during the evolution

process. The two primary variation operators employed in HyperNEAT are crossover and mutation.

As the algorithm progresses, genomes and the neural networks they generate may increase in complexity. The discovery of symmetries and regularities begins, though they do not always pertain fundamentally to the task. For instance, in earlier generations, the substrate generated by CPPN produces an output that results in poor creature behavior, inhibiting movements towards the target. However, as the evolutionary process continues, it uncovers the essential regularity necessary for genomes to achieve optimum ftness for the task.

2.5 Excessive elitism method

In the evolutionary process in NEAT, various mechanisms have been incorporated to strike a balance between ftness and population diversity. These mechanisms include gene tracking, speciation, and incremental complexifcation of the network. NEAT defnes two levels of elitism: species elitism and reproduction elitism. Our focus is on modifying the latter, which proposes that the most-ft individuals in each species will be preserved as-is from one generation to the next.

As seen in Fig. [6,](#page-4-0) we modifed the standard elitism in NEAT, originally defned as the process where the best-ft *M* individuals in each species are preserved as-is from one generation to the next (Fig. [6](#page-4-0) left). In the case of excessive elitism, the best-ft *M* individuals become elite regardless of their species within the whole population of *N* (Fig. [6](#page-4-0) right), under the assumption of an excessive elitism, *M*. Their genotypes are carried forward to the succeeding generation, with the evaluated ftness values also passed on and used as their ftness value instead of re-evaluating the best-ft individuals in subsequent generations. As such, only the non-elite individuals (*N*-*M*) generated through selection

Fig. 6 A comparative analysis between the excessive elitism (EE) method and the standard elitism in NEAT

and genetic operations require evaluation at each generation. In the EE method, concepts of speciation and ftness sharing are turned off, thereby eliminating the species concept during evolution. We implement a roulette wheel selection method to stochastically select parents for reproduction, with being proportional to the ftness within the population. This approach offers a simpler alternative that leads to a simple population structure. More specifcally, the excessive elitism (EE) method could afect the algorithm by giving priority to preserving the best-ft individuals within a large portion of the population for the purpose of exploration. The large population capacity (*N*) in EE enhances the diversity, signifcantly improving the population's ftness. Additionally, replacing only a few individuals with offspring in each generation contributes to lower computational costs.

This substantially reduces evaluation costs for each generation, especially if the number of elite individuals (*M*) is large. Furthermore, we anticipate that maintaining high diversity in a large population can prevent the population from being stuck in local optima, though the rate of evolution may decelerate due to fewer individuals evolving (*N*-*M*). We adopted the following process: All *N* individuals were placed in a 3D simulation feld; however, only the non-elite individuals (*N-M*) underwent development and movement through their lifetime for ftness evaluation. In contrast, the elite individuals (*M*) remained static, undergoing no development or movement. Each individual was evaluated in the initial generation. Related codes for the model can be found in the supplementary materials.

3 Experiments and results

We conducted the evolutionary experiments of artifcial creatures by implementing HyperNEAT algorithm, utilizing the PUREPLES pure python library. Several key parameters from the default confguration were modifed to improve our model's efficiency in evolving the artificial creatures adapted to our specifc evolutionary model. The contributing parameters are presented in Table [1.](#page-5-0)

We have set the ftness criterion to not exceed a maximum threshold of 10,000. This precaution ensures that the evolutionary process does not terminate before completing a generation's time step. It also serves to maintain the continuity of evolution, irrespective of whether certain individuals have already attained the ftness of 60. The 'reset on extinction' is set to false to avoid creating a completely random population should total extinction occur. The number of species that will be protected from stagnation is set to 1 (species elitism) to prevent total extinction. For more details see reference [\[8](#page-10-7)].

We conducted four evolutionary experiments, each with distinct yet comparable parameters. We applied the

Table 1 Modified parameters in NEAT's configuration	Configuration	Parameters
	Fitness criterion	Max
	Fitness threshold	10,000
	Reset on extinction	False
	Species elitism	
	Elitism (default)	2

Table 2 Configuration of *N* and *M* for NEAT $(2, 20)$, EE $(2, 20)$, EE (82, 100), and EE (18, 20)

Excessive Elite method in the cases of EE (2, 20), EE (82, 100), and EE (18, 20). Case NEAT (2, 20) was conducted to demonstrate the evolutionary process using the existing NEAT selection method.

Table [2](#page-5-1) presents the parameters of population (*N*), elite size (*M*), and the number of evaluating individuals (*N*-*M*) in each specifc case. For each case, we conducted 20 trials to obtain the most precise average results for ftness and block numbers. As indicated in Table [2,](#page-5-1) to compare the advantages of EE and NEAT, we executed NEAT (2, 20) with speciation and fitness sharing enabled as in Fig. [6.](#page-4-0) We assumed EE $(2, 20)$ as a baseline case of EE which featured a smaller population size $(N=20)$ and smaller elite size $(M=2)$. This configuration is typically used in evolutionary experiments when evaluation costs need to be minimized, and it resulted in average ftness to reach local optima (explained later). To overcome the problem, we increased the population size to $N = 100$ and added an excessive elite size of $M = 82$ in the case of EE (82, 100), keeping the number of evaluating individuals consistent at *N*−*M* = 18. We expect that there might be an improvement in the ftness increase. Finally, we conducted EE (18, 20) to demonstrate the use of excessive elitism, where $M = 18$, while maintaining the smaller population size as in EE $(2, 1)$ 20) case. We expected that this case would also lead to an increased average ftness.

We conducted each experiment over 400 generations to standardize our fndings and utilize them for further comparisons in the subsequent sections. We ran the simulation using AMD Ryzen 9 5950X 16-Core Processor and 128 GB memory PC.

The overall results are summarized in Fig. [7.](#page-6-0) The graphs in Fig. [7](#page-6-0)(i) showed the evolution of the average fitness for each trial (gray), and their average (blue) and best (orange) for each case. Figure [7\(](#page-6-0)ii) represents the corresponding evolution of the number of blocks within individuals at the end of their lifetime. The standard deviation was obtained as the objective measure for the diversity of morphology in the population (Fig. [7\(](#page-6-0)ii)). For each trial, we determined the standard deviation in block numbers among the individuals within the population. We then computed and plotted the average of these standard deviations across 20 trials. Figure $7(iii)$ shows the trajectory of evaluated individuals in the last generation of the best trial. For a balanced comparison of ftness increase among the cases, we adjusted the *x*-axis scale based on the evaluation cost, calculated as computation time per generation.

The data presented demonstrate a signifcant increase in ftness when EE was implemented, almost reaching the maximum value and successfully reaching the target ((EE (82, 100) and EE (18, 20)). Conversely, ftness tended to converge towards an intermediate value in cases with a smaller elite size such as EE (2, 20). The total number of blocks showed a tendency to frst increase more than around 10 before evolving to smaller values. In the following sections, we will conduct a comprehensive comparison of these cases.

3.1.1 Evolution of artifcial creatures with the original NEAT algorithm (NEAT (2, 20))

We implemented the NEAT algorithm with speciation and ftness sharing enabled in the case of NEAT (2,20) by setting the minimum species size to 2 and the compatibility threshold to 0.03.

Fig. 7 Comparative results for NEAT (2, 20), EE (2, 20), EE (82, 100), and EE (18, 20). Sub-fgure (**i**) portrays the evolution of ftness across 20 trials (with gray indicating the best ftness in each trial, blue signifying the average of average ftness in each trial, and orange representing the average of best ftness in each trial). Sub-fgure (**ii**)

presents the number of blocks (blue for average, green for best). Subfgure (**iii**) illustrates the trajectory of individuals in the last generation of the best trial. Sub-fgure (**iv**) highlights the speciation process during evolution with NEAT, with each color symbolic of a unique species within a population

As shown in Fig. [7](#page-6-0) (NEAT (2, 20)-i), most trials showed successful evolution of the population and most creatures were able to reach the target. The increase in ftness was comparable to the evolution with EE (82, 100) depicted in (Fig. [7](#page-6-0) (EE $(82, 100)$ —i)). Both cases demonstrated high morphological diversity, as can be seen in Fig. [7](#page-6-0) (NEAT (2, 20—ii) and Fig. [7](#page-6-0) (EE $(82, 100)$ —ii), with the best morphology adapting to approximately 3–5 blocks. The evaluation costs for EE (82, 100) and NEAT (2, 20) were comparable, at 121 s/gen and 82 s/gen respectively.

We can conclude that NEAT enabled the population to keep diversity with enabling speciation and ftness sharing, which is expected to contribute to the performance. While EE method had realized equivalent diversity in a simple single population.

3.1.2 Efects of the excessive elite size in increased population size (EE (2, 20) and EE (82, 100))

We conducted a comparative analysis between EE (2, 20) with EE (82, 100). In this comparison, EE (2, 20) served as the baseline experiment of EE, which had previously served as the solution to reduce the evaluation cost by proposing a smaller population size (N), yet this approach caused premature convergence. To overcome this limitation, we increased the population size to $N=100$ and applied an excessive elite size of $M=82$ into EE (82, 100). In this case, we maintained the same evaluating individual size (*N*−*M*=18), allowing us to compare the changes in the performance of the evolved population.

As depicted in (Fig. [7](#page-6-0) (EE $(2, 20)$ —i), numerous trials showed that the ftness faced premature convergence and stuck in the local optima of around 20. The evaluation cost was 117 s/gen. In contrast, EE (82, 100) showed an increase in fitness (Fig. 7 (EE—i)), with fitness reaching the maximum value (60) in many cases. The evaluation cost was comparable at 121 s/gen with that in EE (2, 20). This slight diference in evaluation cost between 121 s/gen and 117 s/ gen might be due to the computational cost for simulating the whole physical feld regardless of the number of evaluating individuals.

Despite the size of the evaluating individuals being the same for both cases $(N-M=18)$, as seen in Fig. [7](#page-6-0) (EE (82, 100)—ii), case EE (82, 100) led to increased morphological diversity as demonstrated by the standard deviation. The increased diversity signifcantly contributed to a greater increase in the average fitness. It further affected the course of morphological evolution, resulting in creatures in EE (2, 20) that tended to obtain around 10–15 blocks, as opposed to the fewer average of 3–5 blocks found in creatures in EE $(82, 100)$, as highlighted in Fig. [7](#page-6-0) (EE $(2, 20)$ —ii) and (EE)—ii). Hence, the larger diversity seen in EE (82, 100) facilitated better morphological and behavioral evolution in the creatures, encouraging them to travel towards the target. Conversely, the lower diversity in EE (2, 20) led to a poor morphological development, hindering creatures from generating forward movements as shown in (Fig. 8 (EE $(2, 20)$)) and (Fig. [8](#page-7-0) (EE (82, 100))).

3.1.3 Efects of the excessive elite size in the small population (EE (2, 20) and EE (18, 20))

We conduct a comparison between EE $(2, 20)$ and EE $(18, 12)$ 20) to assess the positive impact of the proposed elitism

Fig. 8 Illustration of the lifetime of successful individuals during the fnal (400th) generation, featuring examples of morphology throughout each of four recurring events. These examples highlight the mor-

phological development and behavioral cycles in NEAT (2, 20), EE (2, 20), EE (82, 100), and EE (18, 20)

when the population size was relatively small $(N=20)$. The elite size was increased from $M=2$ in EE (2, 20) to $M=18$ in EE (18, 20). As depicted in Fig. [7](#page-6-0) (EE (18, 20)—(i)), the average ftness in EE (18, 20) witnessed a substantial improvement (around 30), which were slightly better than in EE (2, 20) (around 20). Many trials in EE (18, 20) are able to reach maximum ftness, successfully approaching the target object. In addition to that, due to the small number of evaluating individuals (*N*-*M*=2) in EE (18, 20), the average evaluation cost was signifcantly reduced to 68 s/gen.

In conclusion, EE demonstrated efficiency in evolution within both large population (EE (82, 100)) and small population size (EE (18, 20)). However, its efectiveness is more prevalent in larger populations as has been discussed in the previous sections. This certainly signifes that EE represents a fexible evolutionary algorithm in the feld of evolutionary computation.

3.2 Analysis of evolved morphology

Figure [9](#page-8-0) shows the morphological development of best individual, evolving from the frst generation (on the left) to the fnal generation (on the right). The fgures were collected from snapshots of best individuals during the last generation in each case. The distinct forms of the creatures from each case demonstrated that varying parameters conditions can induce evolutionary processes resulting in differing morphological developments, which leads to diverse

behavioral patterns. Videos illustrating the behaviors and morphological developments of the creatures in each case during the fnal generation, as demonstrated in Fig. [8](#page-7-0), are available in the supplementary materials.

In conclusion, the population tended to obtain lower performance when individuals were composed of a larger average number of blocks, which might have hindered the movements. Conversely, higher performance was achieved when individuals were comprised of fewer number of blocks, potentially encouraging better freedom of movement to reach the target.

It should be noted that the average number of blocks once increased (around 10–15) in early generations across all cases. This is attributable to the NEAT algorithm, where the initial genes were randomly generated, creating a uniform population of simple networks. This likely resulted in a larger average number of blocks in these early generations. We expect that, at frst, individuals with more blocks had dominated the population due to an evolutionary tendency. However, these large-sized individuals were too complicated to evolve further and obtain more adaptable morphological structures, leading to premature convergence in cases with smaller elite size. On the other hand, excessive elite sizes maintained diversity within the population which enabled individuals with fewer blocks to survive. This also offered them an opportunity to evolve their morphology and behavior to become simpler and more adaptable.

Generations		0th gen \rightarrow 100th gen \rightarrow 200th gen \rightarrow 300th gen \rightarrow 400th gen	
NEAT(2, 20)			
EE(2, 20)			
EE(82, 100)			
EE(18, 20)			

Fig. 9 Observations of morphological transformations and behavioral patterns that the most successful individuals adopt throughout various generations in NEAT (2, 20), EE (2 20), EE (82, 100), and EE (18, 20)

Table 3 Confguration of *N* and *M* for all cases

Cases EE (M, N)	Popula- tion size, N		Elite size, M Evaluating individuals, $(N-M)$
EE(2, 20)	20	2	18
EE(22, 40)	40	22	
EE(42, 60)	60	42	
EE(62, 80)	80	62	
EE (82, 100)	100	82	
EE (102, 120)	120	102	

3.3 Finding optimal values of elite size (*M***) and population size (***N***) to successfully facilitate the evolution**

As deduced from the results of the cases conducted in 3.1, it has been determined that the evolution of creatures is most efficient in case $EE(82, 100)$ for larger population sizes and EE (18, 20) for smaller population sizes. However, it is still uncertain what the optimal values for the elite size, *M*, and the population size, *N*, would be for the best adaptation to the current environment. Therefore, we proceeded to conduct experiments using various confgurations of *N* and *M*, as shown in Table [3.](#page-9-0)

We maintained the value of evaluating individual (*N*-*M*) at 18 for all cases, based on the best case in 3.1 and 3.2 (EE (82, 100)). The values that changes were the population size (N) and elite size (M) to see how it affects the evolution of creatures. The experiments were carried out for 200 generations for all cases and repeated to 20 trials for each case. The results of average best ftness and average computational time were calculated from the average of 20 trials in each case. These were then plotted against the population size (*N*), as shown in Fig. [10.](#page-9-1) The error bars present on the average best ftness depict the maximum standard deviation of the average number of blocks in the population. This serves as a representation of the morphological diversity present in each case. This arrangement allows for a clearer observation of the optimal confgurations that most efectively evolved the creatures.

Analyzing the results illustrated in Fig. [10](#page-9-1), we observed that the average computational time ranged from 117 to 125 s/gen when the evaluating individuals (*N* − *M*) remained consistently at 18. The average best ftness portrayed a slight improvement, from approximately 25 in (EE (2, 20)) to roughly 30 to 35 in (EE (22, 40)), and this value then remained constant for EE (42, 60). On the contrary, the average time experienced a slight increase from 117 s/ gen to approximately 118 s/gen in the case of EE (42,60). Starting from EE (42, 60), we observed a vague increase in the average time, between 120 and 125 s/gen in the case

20 40 60 80 100 120 140 20 40 60 80 100 120 Average time, Average best-fitness, Elite size (M), against Population size (N) Elite size, M Average time (sec/gen) Average best-fitness EE(2, 20) EE(22, 40) EE(42, 60) EE(62, 80) EE(82, 100) EE(102, 120)

Population size, N

Fig. 10 A depiction of the correlation between the average evaluation time, average best ftness, and elite size (*M*) with respect to population size (*N*). The error bars present on the average best ftness are demonstrative of the standard deviation, representing the morphological diversity in each case

of EE (82, 100). However, a signifcant improvement was noticeable in the average best ftness in the EE (62,80) case, with a slight increase also seen for EE (82,100).

Referring to results in Fig. [10,](#page-9-1) larger population size cases often result in larger diversity of morphology as represented by the standard deviation on the average ftness in Fig. [10](#page-9-1) (EE (82, 100)), where it shows the largest diversity obtained among other cases. We can see a slight decrease in diversity for smaller population case of EE (62, 80), followed by EE (42, 60), EE (22, 40), and EE (2, 20). Hence, larger morphology diversity was achieved in cases with a larger population, *N*. However, surprisingly, morphological diversity did not continue to improve for case EE (102, 120), although this case has the largest population size among other cases. Due to this, the average ftness slightly decreases to around 40–50. We strongly expect that EE (102, 120) may have an extremely large ratio of elite to evaluating individual size (*M*:*N*), within the population which potentially leads to a decrease in diversity, thus causing premature convergence. This might be due to the evolution of only the lowest-ftness individuals, when the larger portion of the population (102:18) is exempted from evaluation. We believe that the evolution of this case is too slow to improve that it converges to the local solution.

Therefore, we observed that the best average ftness of the evolved population, which was around 50 to 60, was obtained at EE (82, 100) as demonstrated in 3.1. Consequently, the morphological development is as in Fig. [9](#page-8-0) (EE (82, 100)), while the behavior of creatures during fnal generation resembles those in Fig. [8](#page-7-0) (EE (82, 100)). A video showcasing the behavior of creature in EE (82, 100) is also available in the supplementary videos.

4 Conclusions

We proposed the Excessive Elitism (EE) method, based on a novel use of elitism in NEAT. EE method served as a simpler alternative to speciation and ftness sharing in the existing NEAT algorithm, yielding equivalent morphological diversity that prevents premature convergence. We evolved the morphology and behavior of artifcial creatures in a 3D multi-agent environment, with a simple target-approaching task. In the proposed EE method, we assumed that the evaluated ftness of the best-ft individuals would be succeeded and reused, instead of being reevaluated during subsequent ftness evaluations. This in turn contributes to a lower evaluation cost.

As has been demonstrated, the excessive elite size surprisingly improved the average ftness of the population in both small and large population cases. Moreover, the effect of excessive elitism in large population cases signifcantly reduced the evaluation cost due to the reduced number of individuals being evaluated. Nonetheless, there are limitations on the ratio of elite to evaluating individuals that can be implemented in a population to obtain the optimum performance for our model, as proven in results in 3.3. We have demonstrated that the optimum size for elite and evaluating individuals would be similar to what has been shown in EE (82, 100).

Thus, by applying the EE method, we shall use the advantage of a large population size and enhanced diversity to achieve optimum ftness whilst keeping the evaluation cost minimal. Nevertheless, it is worth noting that certain non-adaptive individuals with initial conditions may accidentally achieve high ftness. These types of problems have been discussed in the evolutionary context [[13\]](#page-11-1) and in relation to evolutionary algorithms (EAs) at large [[14](#page-11-0)]. Despite these challenges, the benefts of the EE method signifcantly outbalance such drawbacks. Additionally, EE method bears similarities to quality-diversity approaches like MAP-Elites, which captures diverse high-performing solutions by partitioning the search space [[15](#page-11-2)]. In this study, the EE method demonstrates the simplest means of securing a niche and preserving diversity within the general framework of a simple genetic algorithm.

In future work, we intend to acknowledge whether the source of the shown performance, derived from the complexity in the search space, is related with the asymmetry between morphology and control [[16](#page-11-3), [17](#page-11-4)]. Furthermore, it is crucial to explore to what extent would the excessive elitism method improves the evolution of artifcial creatures, while keeping the limitations minimal and maintaining conditions that would facilitate the evolution of a more complex interactions between artifcial creatures and their environment through the implementation of niche constructing behavior. We planned to incorporate various elements into the evolutionary process. Expectations lie in evolving each creature with an exemption from interactions for simplicity, to exclusively analyze multiple levels of adaptive mechanisms—these include the developmental evolution of creatures, the efects of niche construction, and ecological inheritance.

Acknowledgements This work was accomplished under "The Education Program for Data Scientist in Future Value Creation" and was supported in part by JSPS/MEXT KAKENHI 21K12058.

Data and code availability The supplementary materials for this article can be found at the following: Source code: [https://doi.org/10.5281/](https://doi.org/10.5281/zenodo.10450560) [zenodo.10450560](https://doi.org/10.5281/zenodo.10450560). Supplementary Videos: Morphological changes and patterns of movements for best individual in EE (2, 20) and EE (82, 100)[—https://fgshare.com/s/7f467ac2e8aa7c40f21d](https://figshare.com/s/7f467ac2e8aa7c40f21d)

References

- 1. Schmitt F, Piccin O, Barbé L, Bayle B (2018) Soft robots manufacturing: a review. Front Robot AI. [https://doi.org/10.3389/frobt.](https://doi.org/10.3389/frobt.2018.00084) [2018.00084](https://doi.org/10.3389/frobt.2018.00084)
- 2. Wehner M, Truby RL, Fitzgerald DJ, Mosadegh B, Whitesides GM, Lewis JA, Wood RJ (2016) An integrated design and fabrication strategy for entirely soft, autonomous robots. Nature 536(7617):451–455. <https://doi.org/10.1038/nature19100>
- 3. Chiba N, Suzuki R, Arita T (2020) Evolution of complex nicheconstructing behaviors and ecological inheritance of adaptive structures in a physically grounded environment. Front Robot AI. <https://doi.org/10.3389/frobt.2020.00045>
- 4. Giráldez R, Díaz-Díaz N, Nepomuceno I, Aguilar-Ruiz JS (2005) An Approach to reduce the cost of evaluation in evolutionary learning. In: Proceedings of computational intelligence and bioinspired systems, 8th international work-conference on artifcial neural networks, pp 804–811
- 5. Stoy K (2022) Efficiency through GPU-based co-evolution of control and pose in evolutionary robotics. In: Proceedings of the ALIFE 2022: the 2022 conference on artifcial life, paper no.: isal_a_00558, 2
- 6. La Cava W, Moore J (2018) Behavioral search drivers and the role of elitism in soft robotics. In: Proceedings of the ALIFE 2018: the 2018 conference on artifcial life, pp 206–213
- 7. Methenitis G, Hennes D, Izzo D, Visser A (2015) Novelty search for soft robotic space exploration. In: Proceedings of the 2015 genetic and evolutionary computation conference, pp 193–200
- 8. Stanley KO, Miikkulainen R (2002) Efficient evolution of neural network topologies. In: Proceedings of the 2002 congress on evolutionary computation. CEC'02, pp 1757–1762
- 9. Stanley KO, D'Ambrosio DB, Gauci J (2009) A hypercube-based encoding for evolving large-scale neural networks. Artif Life 15(2):185–212.<https://doi.org/10.1162/artl.2009.15.2.15202>
- 10. Stanley KO (2007) Compositional pattern producing networks: a novel abstraction of development. Genet Program Evol Mach 8:131–162.<https://doi.org/10.1007/s10710-007-9028-8>
- 11. Jaafar SA, Suzuki R, Komori S, Arita T (2023) How large elite size can facilitate the evolution of artifcial creatures with NEAT. In: Proceedings of the joint symposiums of AROB-ISBC-SWARM 2023, pp 142–147
- 12. Jaafar SA, Suzuki R, Komori S, Arita T (2023) How excessive elitism can facilitate the evolution of morphology and behavior of artifcial creatures with NEAT. In: Proceedings of the ALIFE

2023: the 2023 conference on artifcial life (3-page abstract) (**in press**)

- 13. Montanier JM, Bredeche N (2011) Embedded evolutionary robotics: the (1+1)-restart-online adaptation algorithm. Stud Comput Intell 341:155–169. [https://doi.org/10.1007/978-3-642-18272-3_](https://doi.org/10.1007/978-3-642-18272-3_11) [11](https://doi.org/10.1007/978-3-642-18272-3_11)
- 14. Jin Y, Branke J (2005) Evolutionary optimization in uncertain environments-a survey. IEEE Trans Evol Comput 9(3):303–317. <https://doi.org/10.1109/TEVC.2005.846356>
- 15. Nordmoen J, Veenstra F, Ellefsen KO, Glette K (2021) MAP-Elites enables powerful stepping stones and diversity for modular robotics. Front Robot AI. [https://doi.org/10.3389/frobt.2021.](https://doi.org/10.3389/frobt.2021.639173) [639173](https://doi.org/10.3389/frobt.2021.639173)
- 16. Ito T, Pilat ML, Suzuki R, Arita T (2013) ALife approach for body-behavior predator-prey coevolution: Body frst or behavior frst? Artif Life Robot 18:36–40. [https://doi.org/10.1007/](https://doi.org/10.1007/s10015-013-0096-y) [s10015-013-0096-y](https://doi.org/10.1007/s10015-013-0096-y)

17. Cheney N, Bongard J, Sunspiral V, Lipson H (2016) On the difficulty of co-optimizing morphology and control in evolved virtual creatures. In: Proceedings of the ALIFE 2016, the ffteenth international conference on the synthesis and simulation of living systems, pp 226–233

Publisher's Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Springer Nature or its licensor (e.g. a society or other partner) holds exclusive rights to this article under a publishing agreement with the author(s) or other rightsholder(s); author self-archiving of the accepted manuscript version of this article is solely governed by the terms of such publishing agreement and applicable law.