

Artificial Life Manuscript Submission

Lexicase Selection for Multi-task Evolutionary Robotics

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Abstract. In Evolutionary Robotics, Lexicase selection has proven effective when a single task is broken down into many individual parameterisations. Evolved individuals have generalized across unique configurations of an overarching task. Here, we investigate the ability of Lexicase selection to generalize across multiple tasks, with each task again broken down into many instances. There are three objectives: to determine the feasibility of introducing additional tasks to the existing platform; to investigate any consequential effects of introducing these additional tasks during evolutionary adaptation; and to explore whether the schedule of presentation of the additional tasks over evolutionary time affects the final outcome. To address these aims we use a quadruped animat controlled by a feed-forward neural network with joint-angle, bearing-to-target and spontaneous sinusoidal inputs. Weights in this network are trained using evolution with Lexicase-based parent selection. Simultaneous adaptation in a wall crossing task (labelled *wall-cross*) is explored when one of two different alternative tasks is also present: *turn-and-seek* or *cargo-carry*. Each task is parameterised into 100 distinct variants, and these variants are used as environments for evaluation and selection with Lexicase. We use performance in a single-task *wall-cross* environment as a baseline against which to examine the multi-task configurations. In addition, the objective sampling strategy (the manner in which tasks are presented over evolutionary time) is varied, and so data for treatments implementing uniform sampling, even sampling, or degrees of generational sampling are also presented. The Lexicase mechanism successfully integrates evolution of both *turn-and-seek* and *cargo-carry* with *wall-cross*, though there is a performance penalty compared to single task evolution. The size of the penalty depends on the similarity of the tasks. Complementary tasks (*wall-cross/turn-and-seek*) show better performance than antagonistic tasks (*wall-cross/cargo-carry*). In complementary tasks performance is not affected by the sampling strategy. Where tasks are antagonistic, uniform and even sampling strategies yield significantly better performance than generational sampling. In all cases the generational sampling requires more evaluations and consequently more computational resources. The results indicate that Lexicase is a viable mechanism for multi-task evolution of animat neurocontrollers, though the degree of interference between tasks is a key consideration. The results also support the conclusion that the naive, uniform random sampling strategy is the best choice when considering final task performance, simplicity of implementation, and computational efficiency.

Keywords: multi-objective, many-objective, evolutionary robotics, lexicase selection, transfer learning

1 Introduction

1.1 Motivation

Evolutionary Robotics (ER) uses principles of evolutionary computation to discover behaviours in artificial autonomous systems through continual adaptation of morphologies and controllers (Floreano et al., 2008). The ambitions of ER are aligned with autonomous robotics more generally, aiming to find control architectures for robots that embody a general capability to deal with problems in their worlds (Vargas et al., 2014). In the present work, we consider this general capability to mean two things: that a controller can perform well across variations of a single task (“semi-generalised control”), and that the controller is competent in multiple task domains which do not necessarily overlap. Current progress in ER means that finding controllers with competency both where multiple tasks exist and also where each task entails a multiplicity of individual parameterisations, is a desirable and feasible research objective.

1.2 Lexicase Selection

Alongside fitness metrics, genetic encodings, and other crucial components of evolutionary algorithms, the method of choosing parents for new generations, the “selection operator”, is a key consideration. Early research in evolutionary computing used a single measure of fitness to select parents and the limitations of this approach ultimately highlighted the need to use selection operators that simultaneously consider a number of different dimensions of performance (Mitchell, 1998). Various evolutionary techniques exist to map and explore effective solutions in a multi-objective optimisation problem. Basic normalisation and averaging of a fitness vector producing a single scalar quantity can be applied, though this technique does not generally scale well, tending to discard high-performing solutions where they are weak in another dimension. More advanced algorithms like NSGA-II (Deb et al., 2002) explicitly acknowledge the Pareto front and archive and sort Pareto-dominant solutions. These algorithms select from the optimal front at each iteration, though they do not perform well in many-objective problems (Seada & Deb, 2015), colloquially defined as problems with more than three objectives. Lexicase selection is a novel many-objective selection operator that often selects specialists (Helmuth et al., 2020), i.e. individuals that are effective in a subset of the objective space but not necessarily the best in every objective, and is capable of scaling to at least 200 individual objectives (Moore & Stanton, 2021).

1.3 Our ER problem domain

In previous work (Moore & Stanton, 2017) we explored Lexicase selection in ER where feed-forward neural controllers are optimised to discover quadrupedal walking gaits. Animats must move towards a target while climbing over a wall of varying height. The wall is positioned half way between the animat’s starting point

32 and its target location. We call this task *wall-cross*. Variation in this obstacle constitutes the task param-
33 eterisation (i.e. the semi-generalised control problem). The obstacle can be one of 100 different heights
34 when presented to animats, resulting in a many-objective optimization problem. Desirable gaits are those
35 that successfully negotiate many different heights¹. We observed that Lexicase evolved populations towards
36 high fitness, exploring difficult areas of the solution space whilst maintaining competency on parts of the
37 problem that had already been solved. We compared Lexicase to algorithms that were designed specifically
38 for this problem—those presented in Stanton and Channon (2013)—and found that Lexicase, even without
39 specific parameter tuning, outperformed those algorithms in all cases.

40 **1.4 Expanding the domain to multiple tasks**

41 Species in the natural world have evolved over millions of years to perform well on a variety of arbitrary prob-
42 lems posed by their environments: natural organisms do not evolve in response to a single, clear adaptive
43 pressure in isolation. It is likely that these overlapping pressures are an important driver of the impressive
44 general competences present in nature. Environmental challenges can reinforce each other and have the
45 potential to select strongly for common adaptations that form the basic building blocks for more advanced
46 adaptive responses. Combinations of these adaptations confer specific capabilities and ultimately respond
47 to particular existential struggles encountered by evolving species.

48 With this in mind, the motivation of the present paper is to explore the performance of Lexicase selection
49 in the ER environment outlined above while expanding the range of objectives to include a second task.
50 Alongside the wall crossing task we introduce and explore two new problems: *turn-and-see*, and *cargo-*
51 *carry*.

52 In *turn-and-see*, the wall is removed and the target position is parameterised into one of 100 configurations.
53 Each configuration translates to a placement of the target on a 180 degree arc centered in front of the animat
54 with performance again measured by the proximity to this target.

55 Figure 1 depicts an animat in each of the three task environments. In the *cargo-carry* task, animats are
56 instantiated with a weighted box on their torso. Successful animats in this task have gaits that maintain a
57 relatively stable torso and thus are able to carry the cargo without it falling to the ground. Here, performance
58 is measured by time. Animats accrue fitness until the box falls from the torso to the floor, at which point the
59 simulation is terminated. The task is parameterised by cargo mass: each of the 100 environments of this
60 task simulates a different weight for the animat to carry.

¹It is important to note that in this configuration, animats do not have information about the wall height. Their gaits are blind and driven only by proprioceptive, directional, vestibular and spontaneous cyclical input signals.

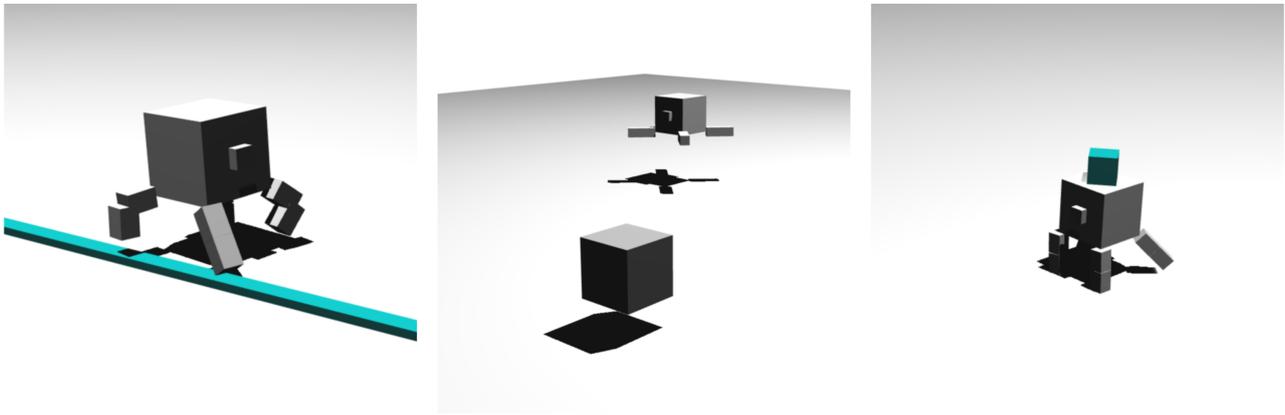


Figure 1: Examples of animat evaluation in the three task environments: *wall-cross* (left); *turn-and-seek* (centre); *cargo-carry* (right). Wall-cross and turn-and-seek both have a target that the animat is navigating toward. This can be seen in the turn-and-seek task as the cube in the foreground.

1.5 Objectives

In the present study there are three main research objectives. First, we wish to determine the feasibility of introducing the additional tasks outlined above to the existing platform. This is in terms of the practicality of integrating them into the algorithm and also to highlight any problems that can occur with the approach.

The second aim is to investigate any consequential effects of introducing these tasks during evolutionary adaptation. These consequences could include reduced performance in one or more tasks or, conceivably, positive synergistic effects if species discover broad, underlying competencies supporting multiple tasks which are threshold discoveries that open domains of even higher fitness.

Third, we aim to explore whether the schedule of presentation of the additional tasks over evolutionary time affects the final outcome. We consider whether a) structuring the presentation of tasks on a generation-by-generation basis, or b) enforcing a certain amount of time spent on each task, has effects on the final outcome of the algorithm in terms of performance of evolved species in specific tasks as well as to the overall performance across the complete problem space.

Broadly, we aim to present a first attempt to show the utility of Lexicase selection in multi-task ER problems. We provide a comparative analysis, contrasting the outcomes of evolution in the multi-task environments with those of single-task populations, a discussion of these results in terms of the interactions between the tasks and subtasks involved, and suggestions for future research directions and expansion of these ideas.

2 Background and Related Work

Evolved robot controllers have proven effective in legged locomotion (Baydin, 2012; Clune et al., 2009; Nolfi & Floreano, 2000) including transferring evolved controllers to physical systems (Koos et al., 2010; Ruud et al., 2016) with fitness often based on the distance traveled in a fixed amount of time. Increasingly, secondary

82 considerations like damage mitigation, and generalizability of behaviours (Pinville et al., 2011) encourage
83 the use of multi-objective algorithms considering multiple performance metrics. Biological observations
84 further enhance systems by bringing in objectives related to efficiency of locomotion (Moore & McKinley,
85 2016). Subsumption architectures (Brooks, 1986; Koza, 1994; Lessin et al., 2013), behavioural diversity ap-
86 proaches (Doncieux & Mouret, 2013) and the combinatorial multi-objective evolutionary algorithm (Huizinga
87 & Clune, 2021) have demonstrated controllers capable of multiple behaviours in one platform. Generalizing
88 controllers spans learning and reacting to environmental contexts across many environments (Lehman et al.,
89 2013), adapting and reconfiguring morphology in response to damage (Kriegman et al., 2019), and exhibiting
90 multiple gaits for one morphology (Cully et al., 2015). Evolving *distinct* behaviours in one controller remains
91 a long-standing goal in ER.

92 Adding generalizability to evolved controllers typically involves moving towards multiple fitness metrics.
93 Multi-objective and many-objective algorithms like Lexicase selection enable scaling fitness objectives into
94 the tens or hundreds of individual objectives. In this study, we expand on earlier investigations (Moore &
95 Stanton, 2017, 2018, 2019, 2020, 2021) by adding new meta-tasks in addition to wall crossing, evaluating
96 the performance of evolved individuals and investigating the performance of Lexicase selection. Adding a
97 second task could lead the evolutionary process to new areas in the search space resulting in higher per-
98 formance (Wagner et al., 2020). Switching between environments can also lead to more effective overall
99 performance across tasks (Canino-Koning et al., 2019; Nahum et al., 2017). Understanding the underly-
100 ing mechanisms that drive Lexicase’s performance, and especially different parameterizations (Hernandez
101 et al., 2022; La Cava et al., 2016) is critical to applying Lexicase effectively. The large search space of
102 many-objective problems can create a computational challenge. Downsampling the number of objectives
103 for consideration during Lexicase selection (Helmuth & Spector, 2020; Hernandez et al., 2019) reduces
104 computational overhead by limiting a selection event to a subset of the objective space. We downsample to
105 10 objectives from a possible 200, consistent with previous wall crossing experiments.

106 **3 Methods**

107 Parameters for the animat, controller, two of three environments, and evolutionary algorithm have been
108 maintained from earlier work (Moore & Stanton, 2021). The software used for these experiments is publicly
109 available and linked at the end of this paper.

110 **Animat Morphology** The quadrupedal animat, shown in Figure 1, has a cube-shaped torso with legs placed
111 at the four lower corners. Each leg has a 2-degree of freedom (DOF) hip and 1-DOF knee. Hips move laterally
112 and vertically allowing the leg to go from straight out from the torso to completely vertical. Knees allow the
113 legs to curl under and towards the torso. Animats get feedback on their position relative to a target through

114 two sensors placed on either side of the torso.

115 **Animat Controller** A feed-forward artificial neural network (ANN) provides control signals for the joints
116 consistent with prior investigations (Moore & Stanton, 2020, 2021). 16 inputs comprising 2 periodic oscil-
117 lating signals, 2 position sensor signals, and 12 for feedback from the joints provide information about the
118 current state of the animat within the simulation. ANNs have one 12 node hidden layer and 12 outputs to
119 control each joint. Genomes are encoded as 336 evolvable weights.

120 **Task Overview** Three primary tasks comprise the objectives in this paper: *wall-cross*, *turn-and-seek*, and
121 *cargo-carry*. Each of the primary tasks is subdivided into 100 unique instances resulting in a total of 300
122 possible objectives. In *wall-cross*, animats are evaluated on their ability to navigate to a target placed in
123 front of the animat beyond a wall. Wall height ranges over 100 values from almost non-existent to the height
124 of the animat's hip, see Figure 2 (left). Figure 2 shows *turn-and-seek*, individuals navigate to a target placed
125 on an arc from the animat's left to right depending on the specific instance. *cargo-carry* places a box of
126 varying density on top of the animat where the box must be carried for as long as possible. Due to the nature
127 of the other two tasks, animats will still attempt to walk toward a target but fitness is not scored based on
128 distance. Rather, fitness is the total time the box is kept aloft, with a possible maximum value of 20 as
129 simulations are conducted for 20 seconds.

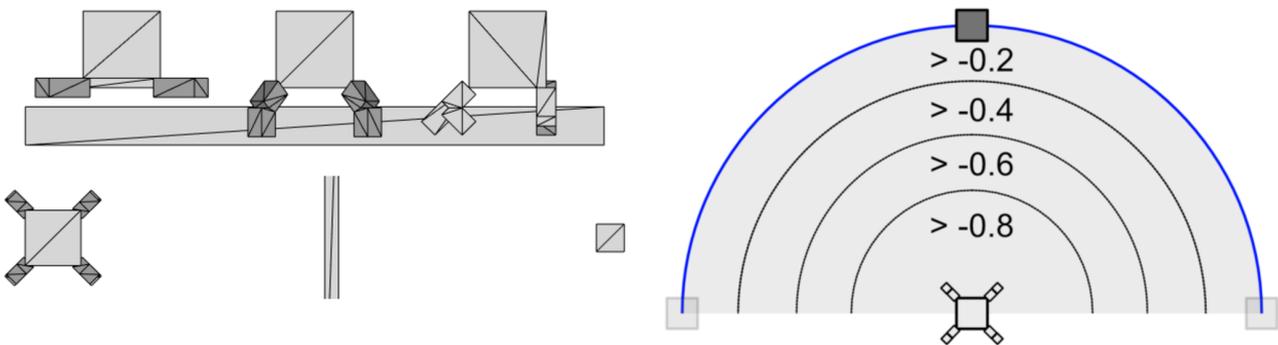


Figure 2: (Left) The maximum wall height in *wall-cross* is set at the animat's hip height if the leg is perpendicular to the ground. Initial position of the animat, wall, and target are shown from a top-down view. (Right) *turn-and-seek* is divided into 100 sub-objectives with the target placed on a semi-circle at 1.8° gradations. Fitness is how close the animat is to the target at a fixed timepoint. (Not to scale.) Radial lines indicate how fitness increases as the animat moves toward the target. Fitness is ultimately a straightline distance to target depending on the specific turn-and-seek environment being evaluated. Figures originally presented in (left) Moore and Stanton, 2020 and (right) Moore and Stanton, 2021.

130 The three tasks together are likely to facilitate some transfer of behaviors as each requires locomotion to
131 be effective. *wall-cross* and *turn-and-seek* are complimentary as effective locomotion and the ability to
132 navigate to a target are behaviors that lead to high performance. However, *cargo-carry* has the potential to
133 be antagonistic to the other two tasks as stable locomotion is favored to elicit high performance in keeping
134 the cargo on top of the animat. This is in opposition to *wall-cross* and *turn-and-seek* which have an emphasis

135 on fast locomotion to reach the target within the simulation time. Prior investigation by Dolson et al., 2018
136 suggests that Lexicase selection may be sensitive to antagonistic objectives.

137 **Evolutionary Algorithm** Downsampled ϵ -Lexicase Selection (Helmuth & Spector, 2020; La Cava et al.,
138 2016) is the evolutionary algorithm used in this study. Individual treatments comprise 20 replicate runs,
139 each with a unique starting seed. 5,000 generations of evolution are run on populations of 50 individuals.
140 A selection event consists of a subsample of five individuals compared on up to 10 objectives from the
141 treatment's objective space. An ϵ - of 10% is applied to comparisons between individuals. As long as an
142 individual is above 90% of the performance of the best individual in the sample for the objective under
143 consideration, it is considered to be tied with the best individual and will move on to performance evaluation
144 on the next objective. If two or more individuals still remain under consideration after all 10 downsampled
145 objectives have been evaluated, a random selection of the remaining individuals in the subsample occurs
146 and a tie event is recorded.

147 **Objective Sampling Strategies** In this paper, we are interested in how multiple tasks, and the interaction
148 between them on evolved controllers, impact performance of Lexicase selection. Treatments are structured
149 based on the tasks they include and how they sample specific task instances in the objective space to best
150 elicit generalized controllers. Three sampling strategies alter how objectives are selected during Lexicase.
151 Note that numbers in the sampling strategy indicate how many tasks are included. The sampling strategies
152 are as follows:

- 153 1. *naive_2t* is the baseline, sampling 10 objectives uniformly per generation across the tasks included in
154 a treatment. Objectives are randomly shuffled so that tasks do not always appear in the same order
155 during Lexicase which would bias the process. Naive sampling is the default behaviour in Lexicase
156 selection.
- 157 2. *even-shuf_2t* samples 5 objectives from each of two tasks per generation. Objectives are then shuffled
158 randomizing ordering during selection preventing one task always appearing first.
- 159 3. *flipN_2t* sample 10 objectives per generation from only one of the two tasks. Once N generations have
160 been selected from this task, a "flip" occurs after which objectives are sampled from the second task
161 resulting in a changing evaluation environment during evolution. For example, with $N = 50$, objec-
162 tives from the first task are selected for 50 generations before the next 50 generations are sampled
163 from the second task. Flipping at every generation (*flip1_2t*) and every 50 generations (*flip50_2t*) are
164 investigated in this study.

165 Table 1 summarises the treatments explored in this study which are variations of the objective sampling
166 strategies discussed above.

Label	Sampling Strategy	Task 1	Task 2
<i>naive_1t</i>	Naive	wall-cross	NA
<i>naive_2t_wt</i>	Naive	wall-cross	turn-and-seek
<i>naive_2t_wt-10000</i>	Naive, 10k gens	wall-cross	turn-and-seek
<i>naive_2t_wc</i>	Naive	wall-cross	cargo-carry
<i>even-shuf_2t_wt</i>	Even Shuffle	wall-cross	turn-and-seek
<i>even-shuf_2t_wc</i>	Even Shuffle	wall-cross	cargo-carry
<i>flip1_2t_wt</i>	Flip every generation	wall-cross	turn-and-seek
<i>flip1_2t_wc</i>	Flip every generation	wall-cross	cargo-carry
<i>flip50_2t_wt</i>	Flip every 50 gens	wall-cross	turn-and-seek
<i>flip50_2t_wc</i>	Flip every 50 gens	wall-cross	cargo-carry

Table 1: Summary of treatments in this study. 20 replicates are undertaken for each treatment. *wt* indicates *wall-turn* environments while *wc* indicates *wall-cargo* environments.

167 4 Results

168 4.1 Wall Crossing Performance One Task vs Two Task

169 In (Moore & Stanton, 2018, 2019, 2020) we investigated characteristics of Lexicase selection in quadrupedal
170 animats evolved for the single task of wall crossing, across 100 unique wall heights (objectives). *naive_1t*
171 provides a baseline of performance replicating results from earlier investigations. Effective wall crossing in
172 these individuals involves both crossing an obstacle of varying height while navigating to the target placed
173 on the opposite side of the wall. Evolved individuals thus have some ability to navigate and locomote which
174 is presumably beneficial in the turning task and also likely detrimental in the cargo task since movement
175 can cause the cargo to fall to the ground. To establish benchmarks of how individuals only evolved for wall
176 crossing versus those evolved to perform wall crossing and a second task, we investigate combinations of
177 two environments as follows.

178 Figure 3 plots the performance of the best individual per replicate for *naive_1t* and *naive_2t_wt* treatments in
179 wall crossing. (The “1t” in *naive_1t* indicates evolution with only one task while “2t” signifies two tasks; “wt”
180 in *naive_2t_wt* represents wall crossing and turn-and-seek as the two evolutionary tasks.) Blue and red dots
181 indicate outliers from the respective treatment’s boxplot distribution both positive or negative. Each box
182 represents the distribution of the best mean performance individual per replicate on the specific objective.
183 Performance for the lower wall heights is nearly identical with most replicates able to reach the target. As
184 wall height increases towards middle heights, individuals evolved only for wall crossing evolve higher per-
185 formance than those evolved for both wall crossing and turning. Wall crossing ability tapers off in both as
186 wall height reaches the upper limits, wherein very specific gaits must evolve to cross these challenging ob-
187 stacles. *naive_1t* does significantly outperform *naive_2t_wt* in the wall crossing task. Statistical significance
188 is determined by a Wilcoxon rank-sum test with Bonferroni correction throughout this paper.

189 Figure 4 plots the performance of the best individual per replicate in the turning task for *naive_1t* and

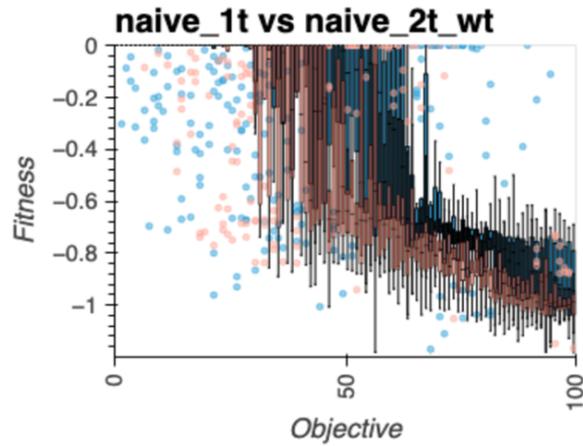


Figure 3: *naive_1t* (blue) and *naive_2t_wt* (red) best individual by mean performance per replicate in wall crossing task. Fitnesses below -0.4 indicate individuals that are unable to cross the wall. Figure adapted from Moore and Stanton, 2021.

190 *naive_2t_wt*. Although not exposed to the turning task during evolution, many of the individuals evolved
 191 only for wall crossing in *naive_1t* are able to navigate to the target when it is placed nearly in front of the
 192 animat (objectives 40-60). As the target is placed further to the left or right, performance tapers off as these
 193 target locations are quite different than those encountered during evolution in wall crossing. *naive_2t_wt*
 194 significantly outperforms *naive_1t* with performance reaching nearly optimal by generation 500. Still, the
 195 ability to navigate to targets placed near the center for individuals in *naive_1t* suggest that the tasks of wall
 196 crossing and turn-and-see may be complementary. That is, behaviours evolved for one task may assist in
 197 solving the other.

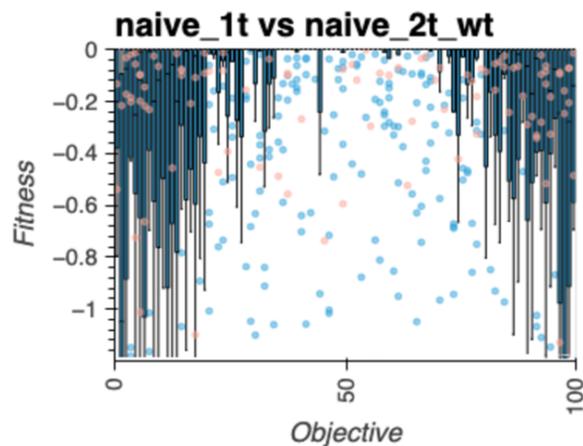


Figure 4: *naive_1t* (blue) and *naive_2t_wt* (red) best individual by mean performance per replicate in turning task. *naive_2t_wt* evolves near perfect turning performance with only a few outliers not evolving full generalization on this task. Figure adapted from Moore and Stanton, 2021.

198 *naive_1t* and *naive_2t_wt* are both evolved for 5,000 generations even though *naive_2t_wt* has 200 objectives
 199 across two tasks. The increase in the number of unique objectives might mean performance differences in
 200 wall crossing are due to fewer selection events occurring for the wall crossing task. We conduct a second two
 201 task naive treatment, *naive_2t_wt-10000*, to see if evolving for 10,000 generations allows for similar per-

202 performance in wall crossing. Figure 5 plots wall crossing performance for the best individual per replicate. No
 203 significant difference in wall crossing performance arises between *naive_1t* and *naive_2t_wt-10000*. Per-
 204 formance between *naive_2t_wt* and *naive_2t_wt-10000* is also not significantly different. Naive selection
 205 strategies are able to evolve similar wall crossing performance, while also incorporating a second task when
 206 allowed to evolve for similar generations relative to the number of tasks. However, our goal with the other
 207 sampling strategies is to assess whether it is possible to evolve generalized two task performance in the
 208 same number of generations as one task performance. Thus, we only evolve for 5,000 generations in the
 209 remaining treatments.

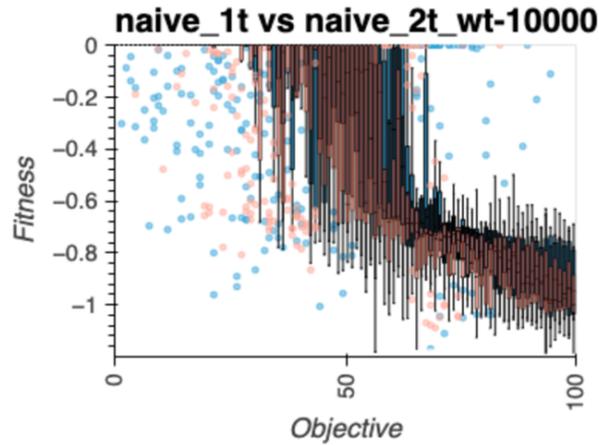


Figure 5: *naive_1t* (blue) and *naive_2t_wt-10000* (red) best mean performance individual per replicate in wall crossing. No significant difference in performance between treatments. Figure originally presented in Moore and Stanton, 2021.

210 Cargo carrying provides another task to evolve individuals alongside wall crossing. Figure 6 plots the wall
 211 crossing performance for individuals evolved only for wall crossing in *naive_1t* and those evolved for wall
 212 crossing and cargo in *naive_2t_wc* ("wc" in *naive_2t_wc* represents wall crossing and cargo carrying as the
 213 two tasks against which agents are evaluated.) Performance in wall crossing of the best individuals evolved
 214 for wall crossing and cargo is significantly lower than that of individuals evolved only for wall crossing after
 215 5,000 generations. Furthermore, Figure 7 plots the cargo carrying performance for the best individuals in
 216 *naive_1t* and *naive_2t_wc*. Performance between the two treatments is significantly different with *naive_2t_wc*
 217 evolving near perfect performance in the cargo carrying task while *naive_1t* individuals have no ability to carry
 218 the cargo no matter the specific cargo density. It takes some time for the box to reach the ground resulting
 219 in fitnesses near 3 for *naive_1t*.

220 For cargo carrying, individuals are not evaluated based on their distance from a target as in wall crossing
 221 and turning. Instead, individuals are measured on their ability to prevent the cargo from falling to the
 222 ground in terms of how many seconds they keep it aloft. The difference in selective pressure results in
 223 there being an antagonistic relationship between wall crossing and cargo carrying. This push-pull between
 224 the tasks invites the hypothesis that, compared to a naive mixing of tasks and objectives, a structured

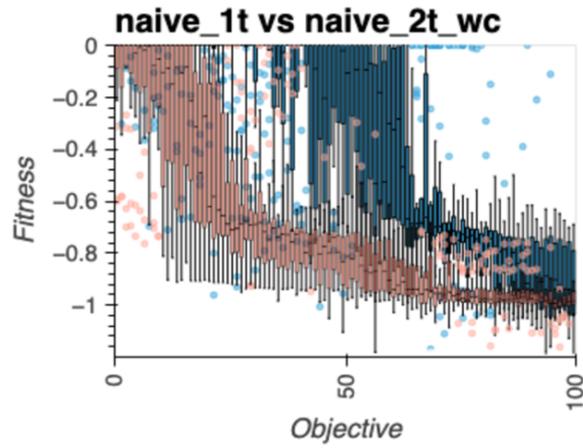


Figure 6: *naive_1t* (blue) and *naive_2t_wc* (red) best mean performance individual per replicate in wall crossing. Performance between the two treatments is significantly different.

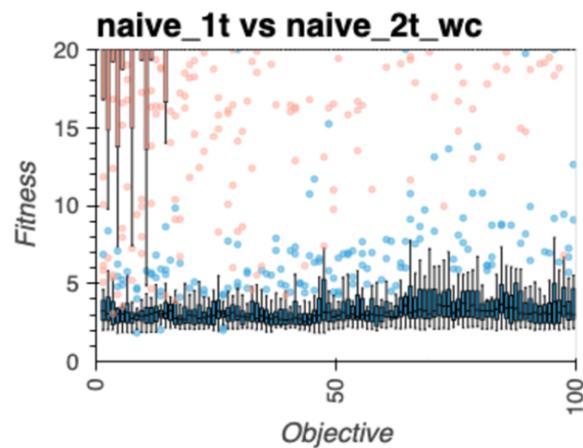


Figure 7: *naive_1t* (blue) and *naive_2t_wc* (red) best mean performance individual per replicate in cargo task. Performance between the two treatments is significantly different. Note the red bars are almost all at 20, which is maximum performance in the cargo carrying task.

225 presentation of objectives over evolutionary time might improve generalised performance on both tasks.
 226 The rationale is that adaptation collapses to one of the two opposing problems when tasks and objectives
 227 are well mixed; by forcing longer adaptive periods on the other task it is possible that sufficient progress is
 228 made to steer populations towards areas of the solution space with potential to achieve high performance
 229 on both problems. This could be due to a differential in the (a posteriori) difficulty of the tasks, or simply a
 230 consequence of the complex dynamics of the adaptive landscape and evolutionary process.

231 4.2 Two Task Complementary Treatments

232 Figure 8 plots the best performing individual per replicate determined by mean performance in wall crossing,
 233 turning, and mean performance for all *wt* treatments. *naive_1t* is also provided as a baseline for comparison.
 234 All two task treatments evolve turning with no significant difference in performance. Wall crossing perfor-
 235 mance is consistent for all two task treatments as well. All *wt* treatments significantly outperform *naive_1t*
 236 in the turning task.

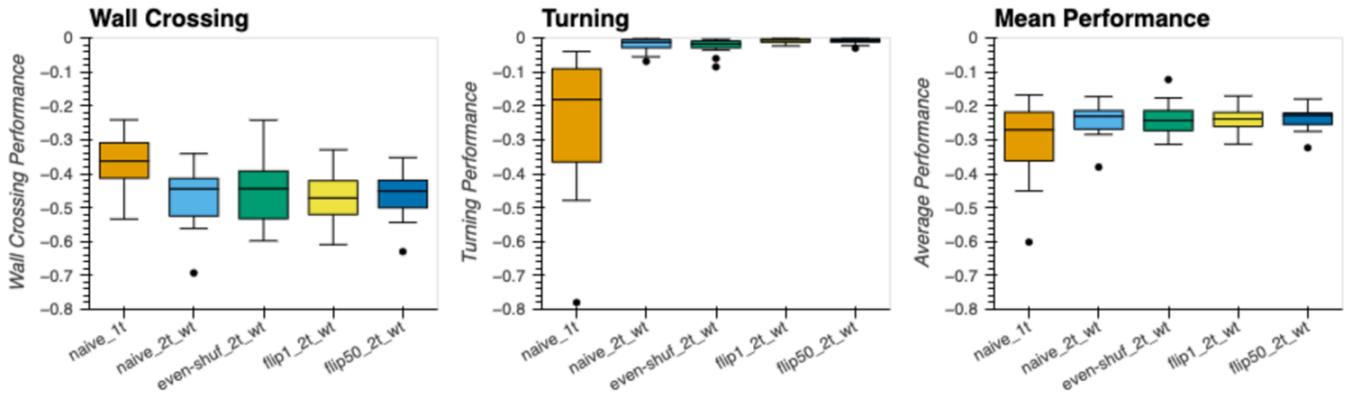


Figure 8: Best individual by mean performance across the two tasks per replicate across treatments run for 5,000 generations. Figure originally presented in Moore and Stanton, 2021.

237 Figure 9 plots the performance per individual objective for the best individual per replicate across the *wt*
 238 treatments. Although performance slightly lags *naive_1t* in all *wt* treatments for wall crossing, generalized
 239 behaviors are evident across *wt* treatments. All *wt* treatments evolve individuals capable of traversing low
 240 wall heights with performance tapering near the middle wall heights. This is consistent with *naive_1t* although
 241 it does have more effective performance on the upper-middle wall heights. Tall walls are difficult to traverse
 242 due to the morphology of the animat and performance tapers off accordingly for all treatments.

243 4.2.1 Lexicase Dynamics

244 Examining the dynamics of Lexicase selection across objective sampling strategies can help elicit differ-
 245 ences that are not apparent from examining performance of evolved individuals. Figure 10 plots the total
 246 number of environments considered during selection per replicate over evolutionary time. A higher num-
 247 ber of selection environments indicates that a sampling strategy had to go farther into the downsampled
 248 objectives to separate individuals during the Lexicase process.

249 Figure 11 plots the number of simulations conducted per replicate. As individuals are filtered out during
 250 selection, the total number of simulations declines. Treatments with fewer individual evaluations might
 251 be good at reducing the subsample of individuals in a selection to one or two performant individuals that
 252 are then tied during the remainder of a Lexicase selection event. Whereas, a high number of selection
 253 environments and subsequently high number of evaluations might mean a treatment is poor at filtering out
 254 individuals throughout selection. Both metrics can indicate the computational efficiency of a treatment.

255 Although performance among the wall and turn sampling strategies are not significantly different, both
 256 *flip* treatments exhibit considerable disparity in both number of environments considered and individual
 257 evaluations between *turn* and *wall* environments while *naive_2t_wt* and *even-shuf_2t_wt* do not. Considering
 258 that the flip treatments sample objectives from only one task per generation it appears that turning is not
 259 as effective as wall crossing at separating individuals out during the selection process.

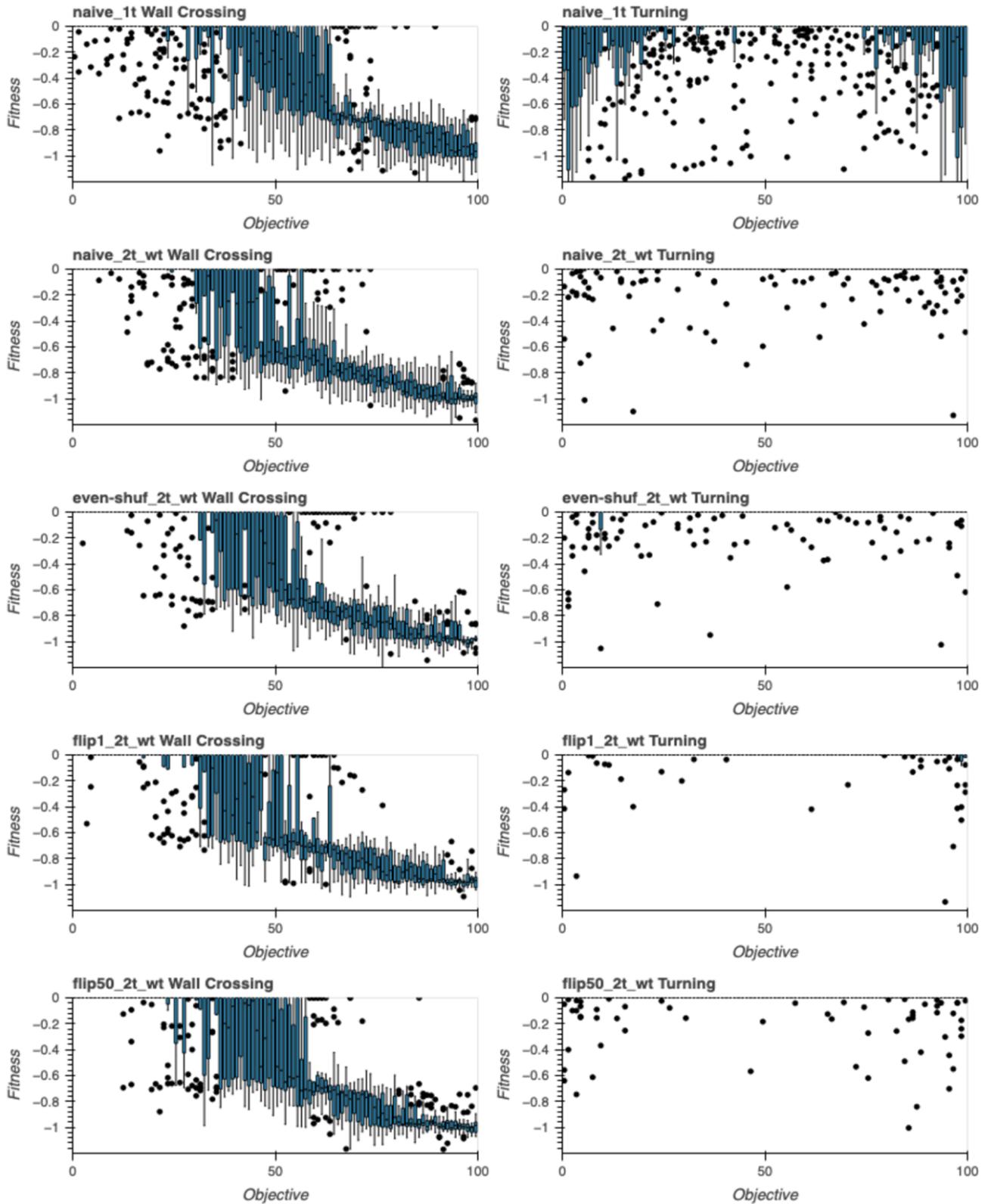


Figure 9: Performance per objective of the best individual by mean performance per replicate across *wt* treatments run for 5,000 generations.

260 *naive_2t_wt* and *even-shuf_2t_wt* do not exhibit a similar difference between the two environments. Both
 261 treatments result in a uniform sampling between environments based on the two figures even though
 262 *naive_2t_wt* does not actively enforce even sampling. Given the differences in Figures 10 and 11 for *flip*
 263 versus *non-flip* treatments, we hypothesize that the turning task is not as effective in filtering individuals as

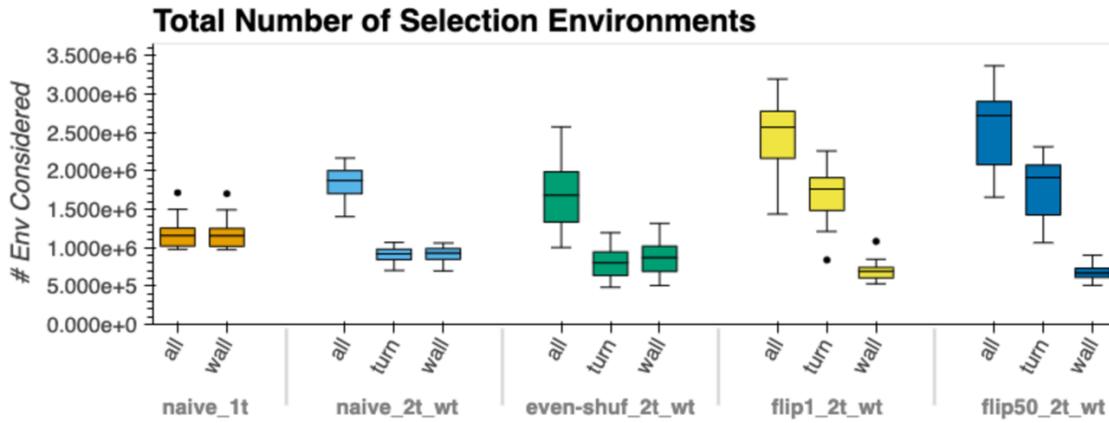


Figure 10: Count of environments considered per replicate. *all* is the sum of wall and turn. Higher numbers suggest more environments were needed per selection event during Lexicase. Figure originally presented in Moore and Stanton, 2021.

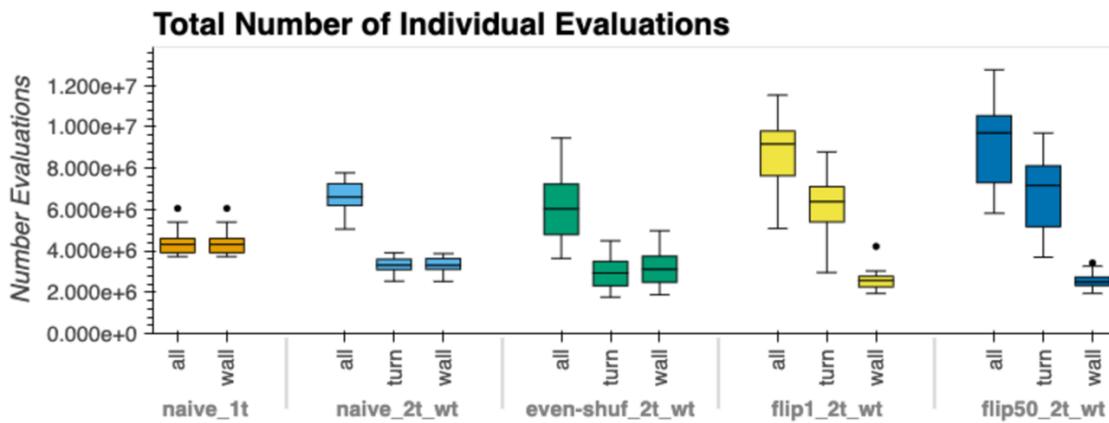


Figure 11: Count of simulations per replicate. *all* is the sum of wall and turn. Simulations represent the significant computational cost. Figure originally presented in Moore and Stanton, 2021.

264 the wall crossing task. Explicitly requiring that only objectives from turning be used for a generation as-in
 265 the *flip* treatments significantly increases computation time for an evolutionary run without a significant
 266 increase in performance in evolved individuals.

267 Figure 12 plots the number of tiebreak events across replicates. Tiebreaks represent a situation where no
 268 individual is determined as better than the others in the sampled selection subset. There is no significant
 269 difference in the number of tiebreaks between *naive_2t_wt* and *even-shuf_2t_wt*, or between *flip1_2t_wt* and
 270 *flip50_2t_wt*. Adding the turning task significantly raises the number of tiebreaks between *naive_1t* and
 271 the two-task treatments while the *flip* strategy further increases tiebreaks. This additional dynamic further
 272 suggests that *flipping* is not an efficient Lexicase selection strategy for these two tasks. Specifically, a high
 273 number of tiebreaks suggests an inability to effectively filter less performant individuals if there is not a
 274 corresponding increase in performance relative to other treatments.

275 Figure 13 plots the percentage of individuals filtered out in a selection event per objective on the left hand
 276 side with the number of times the objective occurs in a replicate on the right hand side. Wall crossing
 277 objectives are numbered 0-99 while turning is 100-199. The figure shows that across treatments, middle

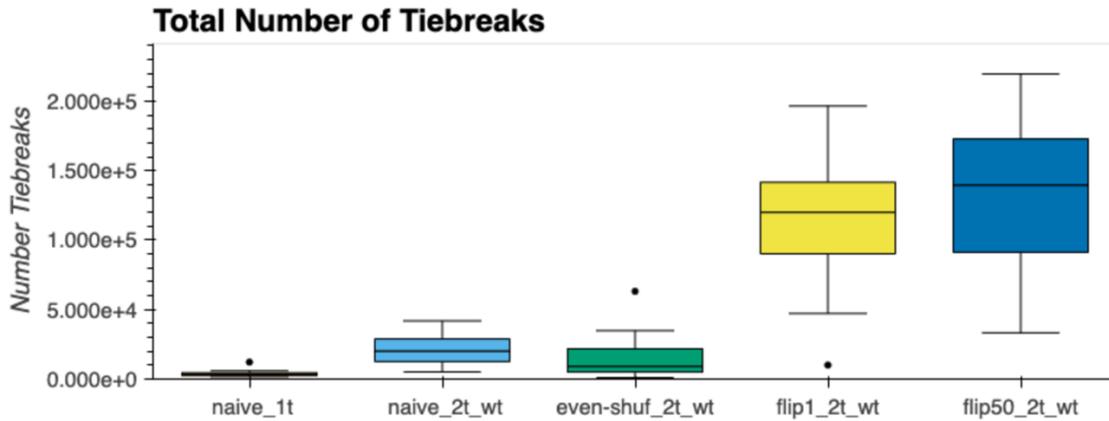


Figure 12: Count of tiebreaks across replicates. Tiebreaks arise when selection cannot isolate one individual during the Lexicase operation. Figure originally presented in Moore and Stanton, 2021.

278 wall heights are the most effective at reducing the number of individuals under consideration by Lexicase
 279 selection. We hypothesize that short walls are relatively easy to cross, assuming an effective locomotion
 280 strategy has evolved, while higher wall heights are also weak selectors as most individuals are incapable of
 281 crossing them. Middle wall heights reach a balance with only the most effective individuals able to cross.
 282 Turning suggests that the far left and far right objectives are the most effective filters while targets placed
 283 in front of the quadrupeds are less effective. Targets placed in front of the quadrupeds share similar sensor
 284 feedback as those of all wall crossing environments likely resulting in some behavioural transfer between the
 285 two tasks. These front facing targets are similar to low wall height objectives in wall crossing which primarily
 286 require an effective locomotion strategy and support the low filtering effectiveness of short wall heights.

287 4.3 Two Task Antagonistic Treatments

288 Figure 14 plots the best performing individual per replicate determined by mean performance in wall cross-
 289 ing, cargo carrying, and mean performance for all *wc* treatments. *naive_1t* is again provided as a base-
 290 line for comparison. All two task treatments evolve cargo carrying ability with no significant difference in
 291 performance, significantly outperforming *naive_1t*. Unlike the turning task, the best evolved wall crossing
 292 individuals from *naive_1t* are not effective at carrying cargo. This suggests that transfer of evolved be-
 293 haviours is reduced between *wc* when compared to results for *wt* in Figure 8. Individuals in *naive_2t_wc*,
 294 *even-shuf_2t_wc*, *flip1_2t_wc*, and *flip50_2t_wc* fail to evolve effective generalized wall crossing behaviour
 295 when compared to their associated treatments in *wt*.

296 Figure 15 plots the performance per individual objective for the best individual per replicate across the *wc*
 297 treatments. In contrast to *wt* treatments, performance is noticeably lower in most wall heights. *naive_2t_wc*
 298 and *even-shuf_2t_wc* are able to evolve some competency on low wall heights while the *flip* treatments show
 299 lower performance. *flip50_2t_wc* is unable to cross any walls while evolving high performance in turning.

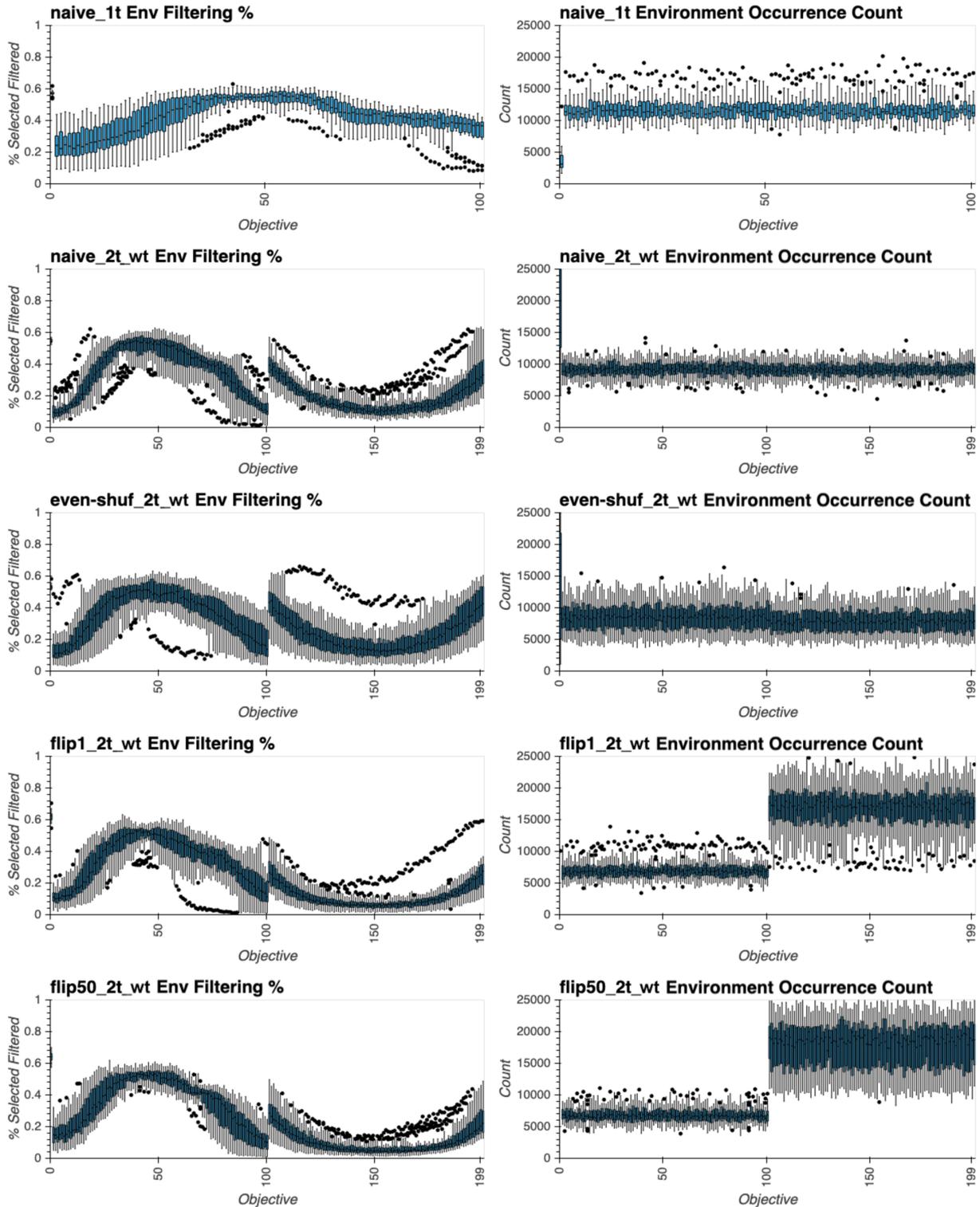


Figure 13: (Left) Filtering percentage during selection distribution across replicates. (Right) Each environment's occurrence count in selection across replicates. Figure originally presented in Moore and Stanton, 2021.

300 4.3.1 Lexicase Dynamics

301 Figures 16 and 17 plot the total number of selection environments and number of individual evaluations per
 302 replicate across treatment for the *wall* and *cargo* tasks. Although the relationship between the two tasks
 303 is different for *wc*, Lexicase dynamics are similar to *wt*. Figure 18 plots the number of tiebreak events per
 304 replicate across treatments. As in the *wt* experiment, the *flip* strategy has significantly higher number of

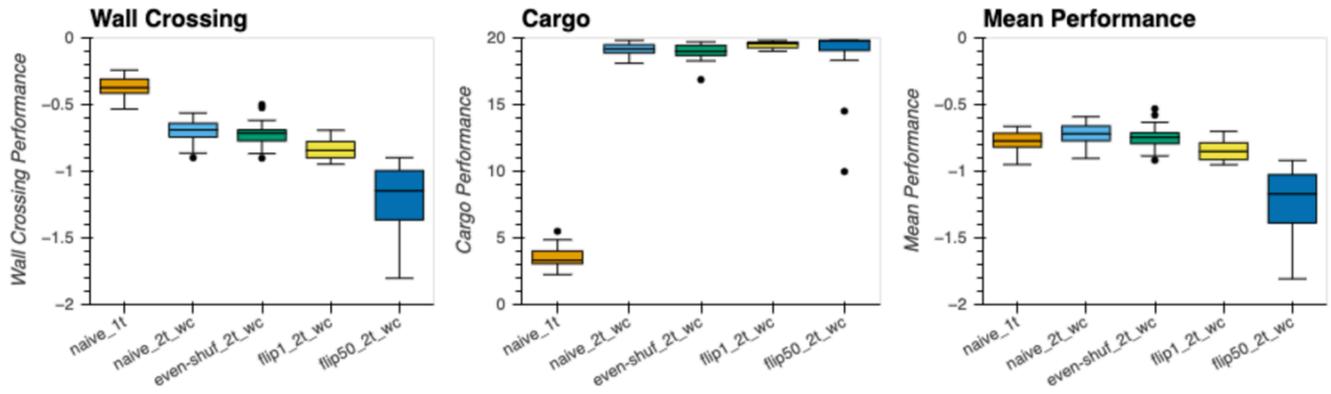


Figure 14: Best individual by mean performance across the two antagonistic tasks per replicate across treatments run for 5,000 generations. Note y-axis for wall crossing is extended to -2 to include *flip50_2t_wc* range.

305 tiebreaks than *naive_2t_wc* and *even-shuf_2t_wc* for two environment treatments.

306 Figure 19 plots (left) the percentage of individuals filtered in a selection event per objective with the (right)
 307 number of times the objective occurs in a replicate. Here, dynamics are different than the *wt* experiments
 308 as the curve of filtering effectiveness for wall crossing objectives shifts towards lower wall heights. This is
 309 especially apparent in the *flip1_2t_wc* and *flip50_2t_wc* treatments. Cargo carrying filtering effectiveness
 310 is also quite low. The high occurrence count of cargo environments in *flip* treatments shown on the right
 311 column of Figure 19, the significantly higher count of tiebreaks versus *naive* and *even shuffling*, and the high
 312 performance in the cargo task together indicate that the general solution to the *cargo-carry* task across
 313 unique box densities is relatively easy to find without behavioural transfer from *wall-cross*.

314 5 Discussion

315 Generalized behaviour is a long-standing goal in robotics. Here, we evaluate Lexicase selection on evolving
 316 quadrupedal animats in two two-task generalization problems. Four objective sampling strategies exhibit
 317 moderate losses in performance in wall crossing compared to the baseline for the *wt* problem with a larger
 318 decrease in the *wc* problem for *flipping* strategies. The small performance differences between the sam-
 319 pling strategies, increases in computational effort for *flip* treatments, and no need to configure additional
 320 parameters suggests that a *naive* objective sampling strategy is effective across both two task problems
 321 examined in this study.

322 Our first objective in this study is to assess the feasibility of integrating multiple environments into the
 323 evolutionary process with Lexicase selection. Notably, individuals in *wt* environments demonstrate both
 324 effective wall crossing on the majority of objectives and the ability to turn and seek a target across objectives.
 325 There are synergies between the two tasks that have likely been exploited in evolved individuals as we see
 326 some ability to execute turn-and-see behaviours in *naive_1t* which does not encounter this task during

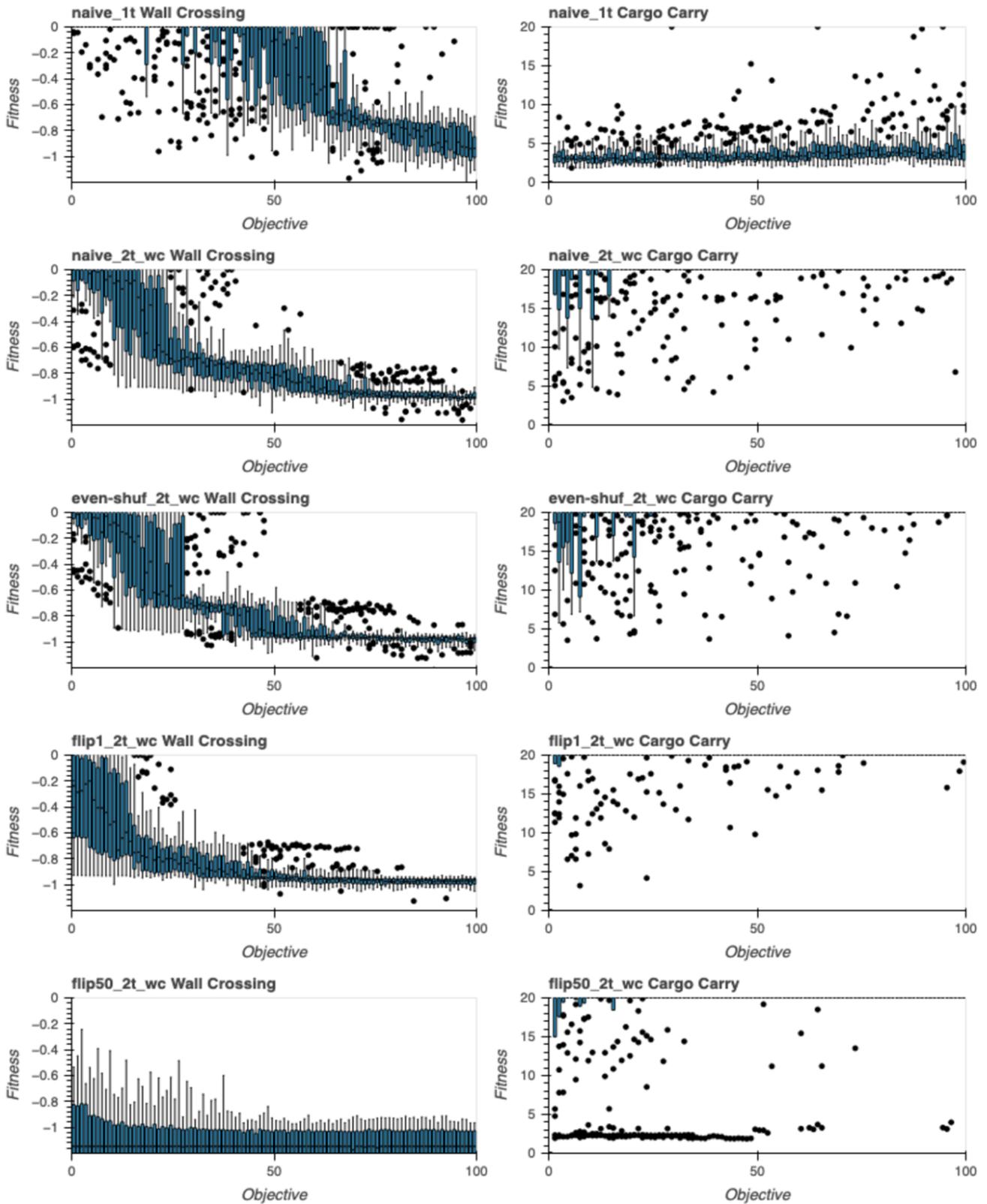


Figure 15: Performance per objective of the best individual by mean performance per replicate across *wc* treatments run for 5,000 generations.

327 evolution. Since both tasks have a target, evolved individuals have their ability to navigate reinforced across
 328 tasks, leading to high performance in both. Individuals evolved in the *wc* environments evolve effective
 329 wall crossing behaviours in lower wall heights only, while they are effective broadly at the cargo carrying
 330 task. Observing evolved individuals in *wc*, carrying a box tends toward small leg movements kept under the

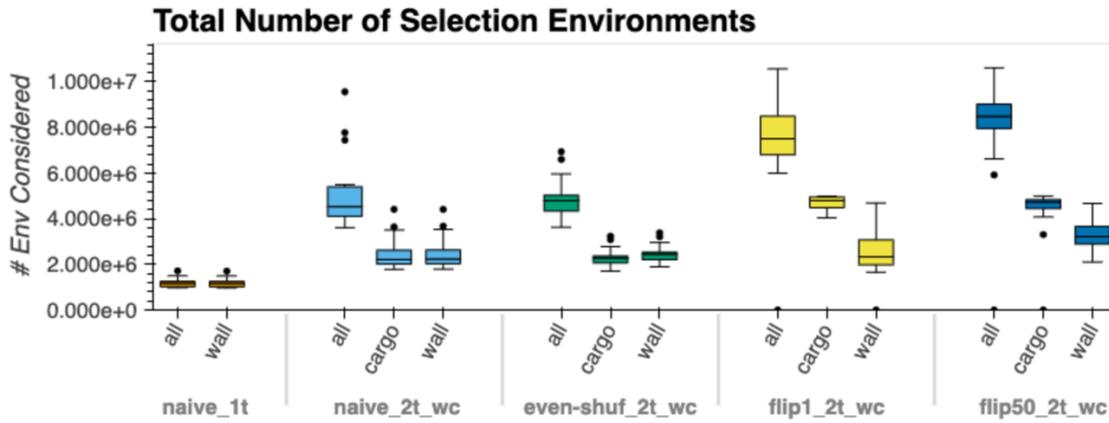


Figure 16: Count of environments considered per replicate. *all* is the sum of wall and cargo. Higher numbers suggest more environments were needed per selection event during Lexicase.

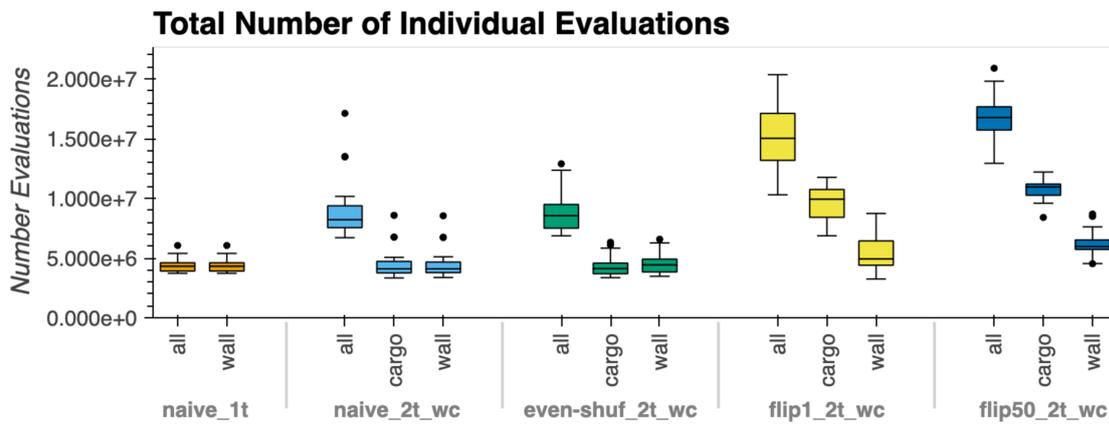


Figure 17: Count of simulations per replicate. *all* is the sum of wall and cargo. Simulations represent the significant computational cost.

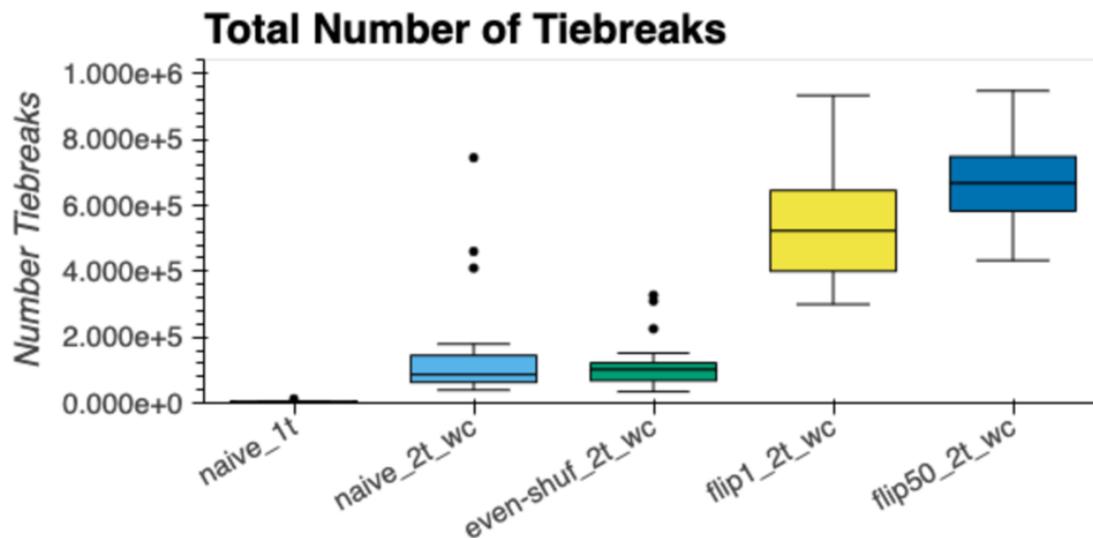


Figure 18: Count of tiebreaks across replicates for antagonistic two tasks.

331 torso maintaining a stable body posture whereas higher wall heights in wall crossing require sweeping leg
 332 movements that reach upwards to step over the obstacle. These conflicting evolutionary pressures likely lead
 333 to the low performance in wall crossing for *wc* treatments and the reduced generalization of individuals in *wc*

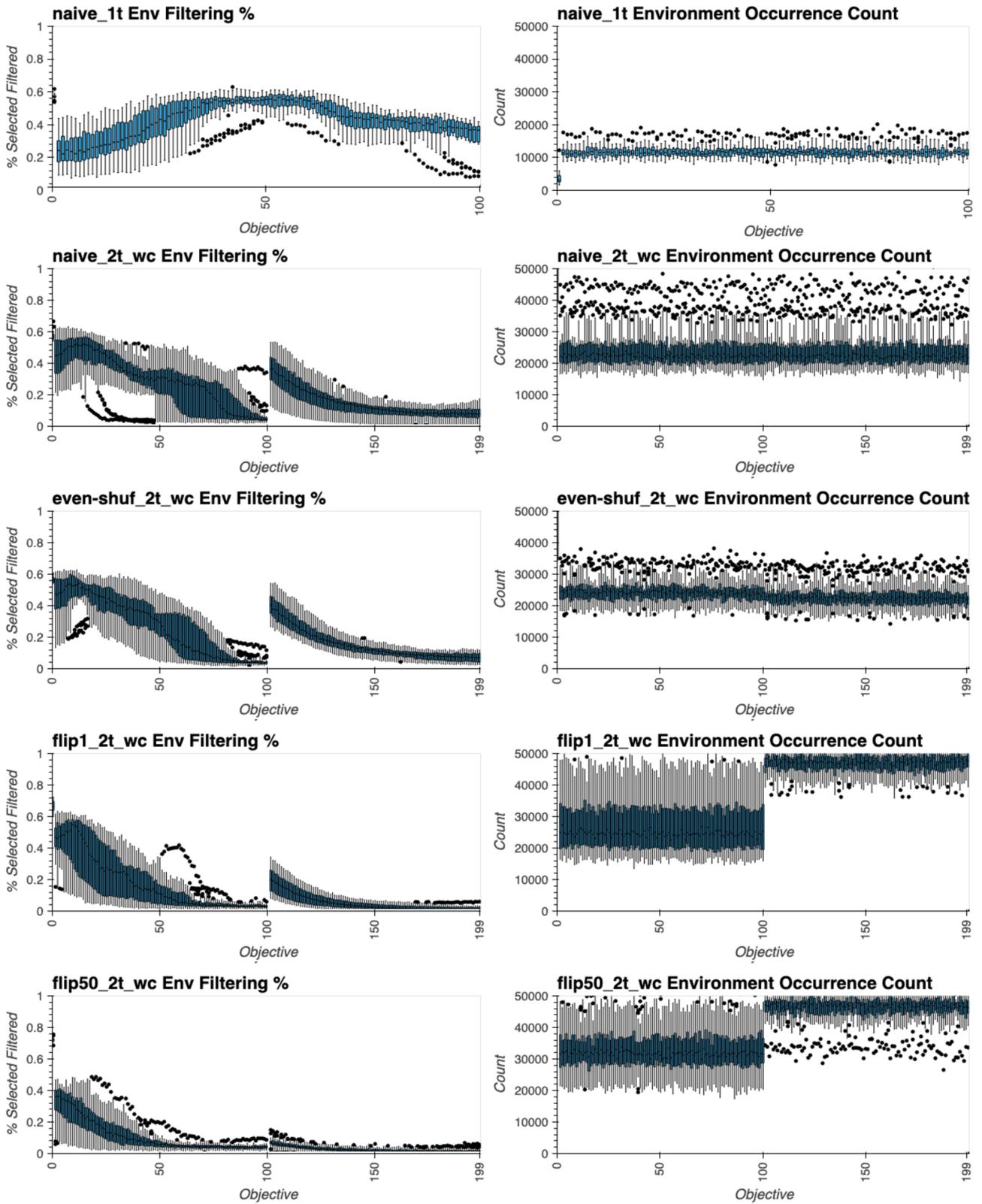


Figure 19: (Left) Filtering percentage during selection distribution across replicates. (Right) Each environment's occurrence count in selection across replicates.

334 versus *wt*. Still, Lexicase selection is able to integrate multiple general tasks effectively in the evolutionary
 335 process. We do observe a decrease in wall crossing performance when adding a second task in both *wt* and
 336 *wc* while keeping the number of generations of evolution stable across the one task baseline and two task
 337 treatments.

338 The second objective in this study is to examine any consequences of introducing the additional tasks,
339 while the third objective explores the schedule of presentation of objectives over evolutionary time. *flipping*
340 strategies are intended to allow the population to specialize on an environment for a specified number of
341 generations. In *flip1_2t_wt* and *flip1_2t_wc*, objectives were sampled from one task per generation with the
342 task flipping every generation. This was intended to prevent one task from dominating selection resulting in
343 poor performance in one task. *flip50_2t_wt* and *flip50_2t_wc* reduced the flipping rate to alternating tasks
344 every 50 generations. Here, individuals would be given a substantial amount of time to specialize on an
345 objective, hypothetically increasing their competency on a task before swapping to the other task. *flipping*
346 strategies appear to be broadly poorer than *naive* or *even shuffling* as their performance is either level
347 with, see Figure 8, or substantially reduced across all environments, see Figure 14, considered in this study.
348 Moreover, when tasks are antagonistic as in the case of *wc*, it appears that *flip50_2t_wc* overspecializes
349 on the cargo task, resulting in a failure to generalize to the wall crossing task. Whereas, *flip1_2t_wc* is able
350 to maintain slightly lower performance than *naive_2t_wc* and *even-shuf_2t_wc* in wall crossing. Against of
351 our intuition that allowing replicates to evolve in one environment for an extended number of generations
352 would improve competency on the tasks, the extra time given to specialize may in fact be detrimental to
353 generalization. This result could be due to either catastrophic forgetting of the second task or evolution of
354 a specialized behaviour for one task that preventing effective behaviours in the second task. In addition,
355 there is a significantly higher number of evaluations and tiebreaks in both two environment combinations
356 when compared to *naive* or *even-shuffling* strategies.

357 **6 Future Work**

358 In future work, we plan to expand the number of tasks while exploring how to better integrate tasks that
359 introduce conflicting selective pressures. Of specific interest will be any changes in performance or charac-
360 teristics of Lexicase selection that might arise in those broader search spaces. We also plan to leverage the
361 filtering efficacy of objectives to explore whether an *adaptive* strategy favoring individual objectives might
362 enhance performance of the algorithm in large search spaces.

363 **7 Acknowledgements**

364 The authors thank William La Cava, Thomas Helmuth, Edward Pantridge, Emily Dolson, and Lee Spector for
365 their conversations about Lexicase selection. We would also like to acknowledge Aston University, Keele
366 University, and Grand Valley State University for supporting this research. The software used for this project
367 has been made available via the public repository linked below².

²<https://github.com/machine-machines/lexicase-alife-journal-2022>

368 **References**

- 369 Baydin, A. G. (2012). Evolution of central pattern generators for the control of a five-link planar bipedal walking
370 mechanism. *Paladyn. Journal of Behavioral Robotics*, 3(1), 45–53.
- 371 Brooks, R. A. (1986). A robust layered control system for a mobile robot. *IEEE Journal of Robotics and Au-*
372 *tomation*, 2(1), 14–23.
- 373 Canino-Koning, R., Wiser, M. J., & Ofria, C. (2019). Fluctuating environments select for short-term phenotypic
374 variation leading to long-term exploration. *PLOS Computational Biology*, 15(4), 1–32. [https://doi.org/](https://doi.org/10.1371/journal.pcbi.1006445)
375 [10.1371/journal.pcbi.1006445](https://doi.org/10.1371/journal.pcbi.1006445)
- 376 Clune, J., Beckmann, B. E., Ofria, C., & Pennock, R. T. (2009). Evolving coordinated quadruped gaits with the
377 HyperNEAT generative encoding, In *Proceedings of the IEEE congress on evolutionary computation*,
378 Trondheim, Norway, IEEE.
- 379 Cully, A., Clune, J., Tarapore, D., & Mouret, J.-B. (2015). Robots that can adapt like animals. *Nature*, 521(7553),
380 503–507. <https://doi.org/10.1038/nature14422>
- 381 Deb, K., Pratap, A., Agarwal, S., & Meyarivan, T. (2002). A fast and elitist multiobjective genetic algorithm:
382 NSGA-II. *IEEE Transactions on Evolutionary Computation*, 6(2), 182–197. [https://doi.org/10.1109/](https://doi.org/10.1109/4235.996017)
383 [4235.996017](https://doi.org/10.1109/4235.996017)
- 384 Dolson, E., Banzhaf, W., & Ofria, C. (2018). Applying ecological principles to genetic programming (W. Banzhaf,
385 R. S. Olson, W. Tozier, & R. Riolo, Eds.). In W. Banzhaf, R. S. Olson, W. Tozier, & R. Riolo (Eds.), *Genetic*
386 *programming theory and practice xv*, Springer International Publishing.
- 387 Doncieux, S., & Mouret, J. B. (2013). Behavioral diversity with multiple behavioral distances, In *Proceedings*
388 *of the 2013 IEEE congress on evolutionary computation*, Cancun, Mexico, IEEE.
- 389 Floreano, D., Husbands, P., & Nolfi, S. (2008). Evolutionary Robotics. In *Handbook of Robotics*. Berlin, Springer
390 Verlag.
- 391 Helmuth, T., Pantridge, E., & Spector, L. (2020). On the importance of specialists for lexibase selection.
392 *Genetic Programming and Evolvable Machines*, 21(3), 349–373. [https://doi.org/10.1007/s10710-](https://doi.org/10.1007/s10710-020-09377-2)
393 [020-09377-2](https://doi.org/10.1007/s10710-020-09377-2)
- 394 Helmuth, T., & Spector, L. (2020). Explaining and exploiting the advantages of down-sampled lexibase se-
395 lection, In *The 2020 conference on artificial life: A hybrid of the European conference on artificial*
396 *life (eCAL) and the international conference on the synthesis and simulation of living systems (alife)*,
397 Montreal, Quebec, Canada, MIT Press. https://doi.org/10.1162/isal_a_00334
- 398 Hernandez, J. G., Lalejini, A., Dolson, E., & Ofria, C. (2019). Random subsampling improves performance in
399 lexibase selection, In *Proceedings of the genetic and evolutionary computation conference compan-*
400 *ion*, Prague, Czech Republic, ACM. <https://doi.org/10.1145/3319619.3326900>

- 401 Hernandez, J. G., Lalejini, A., & Ofria, C. (2022). An exploration of exploration: Measuring the ability of lexicase
402 selection to find obscure pathways to optimality. In W. Banzhaf, L. Trujillo, S. Winkler, & B. Worzel
403 (Eds.), *Genetic programming theory and practice xviii* (pp. 83–107). Singapore, Springer Singapore.
404 https://doi.org/10.1007/978-981-16-8113-4_5
- 405 Huizinga, J., & Clune, J. (2021). Evolving Multimodal Robot Behavior via Many Stepping Stones with the Com-
406 binatorial Multi-Objective Evolutionary Algorithm. *Evolutionary Computation*, [https://direct.mit.edu/evco/article-](https://direct.mit.edu/evco/article-pdf/doi/10.1162/evco_a_00301/1974750/evco_a_00301.pdf)
407 [pdf/doi/10.1162/evco_a_00301/1974750/evco_a_00301.pdf](https://direct.mit.edu/evco/article-pdf/doi/10.1162/evco_a_00301/1974750/evco_a_00301.pdf), 1–34. [https://doi.org/10.1162/evco_](https://doi.org/10.1162/evco_a_00301)
408 [a_00301](https://doi.org/10.1162/evco_a_00301)
- 409 Koos, S., Mouret, J. B., & Doncieux, S. (2010). Crossing the reality gap in evolutionary robotics by promot-
410 ing transferable controllers, In *Proceedings of the 2010 acm genetic and evolutionary computation*
411 *conference*, Portland, Oregon, USA, ACM. <https://doi.org/10.1145/1830483.1830505>
- 412 Koza, J. R. (1994). Evolution of a subsumption architecture that performs a wall following task for an au-
413 tonomous mobile robot, In *Proceedings of the workshop on computational learning theory and nat-*
414 *ural learning systems : Intersections between theory and experiment*, Berkeley, California, USA, MIT
415 Press.
- 416 Kriegman, S., Walker, S., Shah, D. S., Kramer-Bottiglio, R., & Bongard, J. C. (2019). Automated shapeshifting for
417 function recovery in damaged robots (A. Bicchi, H. Kress-Gazit, & S. Hutchinson, Eds.). In A. Bicchi, H.
418 Kress-Gazit, & S. Hutchinson (Eds.), *Robotics: Science and systems xv, university of freiburg, freiburg*
419 *im breisgau, germany, june 22-26, 2019*, University of Freiburg. [https://doi.org/10.15607/RSS.2019.](https://doi.org/10.15607/RSS.2019.XV.028)
420 [XV.028](https://doi.org/10.15607/RSS.2019.XV.028)
- 421 La Cava, W., Spector, L., & Danai, K. (2016). Epsilon-lexicase selection for regression, In *Proceedings of the*
422 *genetic and evolutionary computation conference 2016*, Denver, Colorado, USA, ACM.
- 423 Lehman, J., Risi, S., D'Ambrosio, D., & Stanley, K. O. (2013). Encouraging reactivity to create robust machines.
424 *Adaptive Behavior - Animals, Animats, Software Agents, Robots, Adaptive Systems*, 21(6), 484–500.
425 <https://doi.org/10.1177/1059712313487390>
- 426 Lessin, D., Fussell, D., & Miikkulainen, R. (2013). Open-ended behavioral complexity for evolved virtual crea-
427 tures, In *Proceedings of the 2013 acm genetic and evolutionary computing conference*, Amsterdam,
428 Netherlands, ACM.
- 429 Mitchell, M. (1998). *An introduction to genetic algorithms*. MIT press.
- 430 Moore, J. M., & McKinley, P. K. (2016). A comparison of multiobjective algorithms in evolving quadrupedal
431 gaits (E. Tuci, A. Giagkos, M. Wilson, & J. Hallam, Eds.). In E. Tuci, A. Giagkos, M. Wilson, & J. Hallam
432 (Eds.), *From animals to animats 14: Proceedings of the 14th international conference on simulation*
433 *of adaptive behavior, sab 2016*, Aberystwyth, UK, Springer International Publishing.

- 434 Moore, J. M., & Stanton, A. (2017). Lexicase selection outperforms previous strategies for incremental evolu-
435 tion of virtual creature controllers, In *Proceedings of the 14th european conference on artificial life*,
436 Lyon, France, MIT Press.
- 437 Moore, J. M., & Stanton, A. (2018). Tiebreaks and diversity: Isolating effects in lexicase selection, In *Proceed-*
438 *ings of the 16th international conference on the simulation and synthesis of living systems*, Tokyo,
439 Japan, ACM.
- 440 Moore, J. M., & Stanton, A. (2019). The limits of lexicase selection in an evolutionary robotics task, In *The*
441 *2019 conference on artificial life: A hybrid of the european conference on artificial life (ecal) and the*
442 *international conference on the synthesis and simulation of living systems (alife)*, Newcastle-Upon-
443 Tyne, UK, MIT Press.
- 444 Moore, J. M., & Stanton, A. (2020). When specialists transition to generalists: Evolutionary pressure in lexicase
445 selection, In *Artificial life conference 2020 proceedings*, Online. MIT Press.
- 446 Moore, J. M., & Stanton, A. (2021). Objective sampling strategies for generalized locomotion behavior with
447 lexicase selection, In *Artificial life conference 2021 proceedings*, Online, MIT Press. MIT Press. https://doi.org/10.1162/isaL_a_00398
448
- 449 Nahum, J. R., West, J., Althouse, B. M., Zaman, L., Ofria, C., & Kerr, B. (2017). Improved adaptation in exoge-
450 nously and endogenously changing environments. https://doi.org/10.1162/isaL_a_052
- 451 Nolfi, S., & Floreano, D. (2000). *Evolutionary robotics: The biology, intelligence and technology of self-*
452 *organizing machines*. The MIT Press.
- 453 Pinville, T., Koos, S., Mouret, J.-B., & Doncieux, S. (2011). How to promote generalisation in evolutionary
454 robotics: The progab approach, In *Proceedings of the 13th annual conference on genetic and evo-*
455 *lutionary computation*, Dublin, Ireland, ACM. <https://doi.org/10.1145/2001576.2001612>
- 456 Ruud, E. L., Samuelsen, E., & Glette, K. (2016). Memetic robot control evolution and adaption to reality, In
457 *Proceedings of the 2016 IEEE symposium series on computational intelligence (ssci)*, Athens, Greece,
458 IEEE.
- 459 Seada, H., & Deb, K. (2015). U-nsga-iii: A unified evolutionary optimization procedure for single, multiple, and
460 many objectives: Proof-of-principle results (A. Gaspar-Cunha, C. Henggeler Antunes, & C. C. Coello,
461 Eds.). In A. Gaspar-Cunha, C. Henggeler Antunes, & C. C. Coello (Eds.), *Evolutionary multi-criterion*
462 *optimization*, Cham, Springer International Publishing.
- 463 Stanton, A., & Channon, A. (2013). Heterogeneous complexification strategies robustly outperform homo-
464 geneous strategies for incremental evolution, In *Proceedings of the 12th european conference on*
465 *artificial life*, Taormina, Italy, MIT Press.
- 466 Vargas, P. A., Paolo, E. A. D., Harvey, I., & Husband, P. (Eds.). (2014). *The horizons of evolutionary robotics*.
467 MIT Press.

468 Wagner, A. P., Zaman, L., Dworkin, I., & Ofria, C. (2020). Behavioral strategy chases promote the evolution
469 of prey intelligence*. In W. Banzhaf, B. H. Cheng, K. Deb, K. E. Holekamp, R. E. Lenski, C. Ofria,
470 R. T. Pennock, W. F. Punch, & D. J. Whittaker (Eds.), *Evolution in action: Past, present and future:
471 A festschrift in honor of erik d. goodman* (pp. 225–246). Springer International Publishing. [https:
472 //doi.org/10.1007/978-3-030-39831-6_17](https://doi.org/10.1007/978-3-030-39831-6_17)