Mouse Academy: high-throughput automated training and trial-by-trial behavioral analysis during learning

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ABSTRACT

Progress in understanding how individual animals learn will require high-throughput standardized methods for behavioral training but also advances in the analysis of the resulting behavioral data. In the course of training with multiple trials, an animal may change its behavior abruptly, and capturing such events calls for a trial-by-trial analysis of the animal's strategy. To address this challenge, we developed an integrated platform for automated animal training and analysis of behavioral data. A low-cost and space-efficient apparatus serves to train entire cohorts of mice on a decision-making task under identical conditions. A generalized linear model (GLM) analyzes each animal's performance at single-trial resolution. This model infers the momentary decision-making strategy and can predict the animal's choice on each trial with an accuracy of ~80%. We also assess the animal's detailed trajectories and body poses within the apparatus. Unsupervised analysis of these features revealed unusual trajectories that represent hesitation in the response. This integrated hardware/software platform promises to accelerate the understanding of animal learning.

INTRODUCTION

Learning – the change of neural representation and behavior that results from past experience and the consequences of actions – is important for animals to survive and forms a central topic in neuroscience[1]. Different individuals may apply different strategies to the learning process, reflecting their individual personalities. Indeed, substantial differences in sensory biases, locomotion, motivation, and cognitive competence have been observed in populations of fruit flies[2,3], rodents and primates[4-6]. Thus, it is critical to investigate learning at the individual level.

Rodents, especially the mouse, have become popular experimental animals in studying associative learning and decision-making, because of the wide availability of transgenic resources[7-10]. They can learn to perform complex decision-making tasks that probe cognitive components such as working memory and selective attention[11-13]. However, differences in learning strategies across individuals have rarely been addressed, partly owing to the limitations of data gathering and analysis.

Studying differences among individuals requires training and collecting data from multiple animals in a standardized and high-throughput fashion. The training procedures are often time-

consuming, requiring several days to many weeks[8,9], depending on the task. Although there have
been advances in training automation, existing systems either require an experimenter to move
animals from the home cage to the training apparatus[14-16], or training animals within their own
cages[17-19]. The former introduces additional sources of variability[20,21], and the latter precludes
tasks that require a large training arena. Following data acquisition, the analysis of behavior aims
at understanding the learning process. Present approaches tend to focus on the averaged
performance over many trials[22]. However, changes in behavior may happen at a single trial, and
thus the modeling of behavior should similarly offer a time resolution of single trials to assess
each animal's individual approach to learning.
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To address these challenges, we present Mouse Academy, an integrated platform for automated
training of group-housed mice and analysis of behavioral changes in learning a decision-making
task. We designed hardware that makes use of implanted radio frequency identification (RFID)
chips to identify each mouse, and guides the animal into a behavior training box. Synchronized
video recordings and decision-making sequences are acquired during animal learning. To
analyze the decision-making sequences, we developed an iterative generalized linear model
(GLM). This model makes a prediction of the animal's choice in each trial and gets updated
based on the animal's actual choice. This iterative GLM model achieves a prediction accuracy of
~80%, and also reveals the decision-making strategy of the animal and how it changes over time.
To analyze the animal's behavior during the task in greater detail, we computed the movement
trajectories of each mouse. These trajectories allowed us to perform an unsupervised analysis of
each animal's behavior, and discover individual traits of behavioral learning that were not
apparent from the simple choice sequences.
72
73 **RESULTS**
74 The Mouse Academy platform consists of three components (**Fig. 1**): an automated RFID sorting
75 and animal training system, an iterative GLM to analyze decision-making sequences, and
76 behavior assessment software that analyzes animal trajectories computed from video data.
77
Automated RFID sorting supports individual training programs
We designed the equipment in the following manner (**Fig. 1a**): RFID-tagged mice are grouped in
a common home cage where food and bedding is supplied. The home cage connects to a
behavior training box through a gated tunnel. The gates are controlled by a home-made RFID
animal sorting system[23]: three RFID antennas are placed along the tunnel, with one near the
home cage, one near the training box and one between the two; the motorized gates are placed
between the RFID sensors, separating the tunnel into three compartments. An Arduino
microcontroller integrates information from the RFID readers to open and shut the gates,
allowing only one animal at a time to pass through the tunnel (**Supplementary Figs. 1a, 1c, 1d**
and **Supplementary Videos 1-4**). The behavior box is outfitted with three ports, each of which
contains a photo-transistor to detect snout entry, a solenoid valve to deliver water reward, and a
light emitting diode (LED) to present visual cues. To maintain a controlled environment, the
training box is isolated from the outside by a light- and sound-proof chamber (**Supplementary
Fig. 1b**).
92
Once a mouse enters the training box, a protocol is set up to train the mouse to perform a certain
task. In the experiments reported here, the animal must nose-poke the center port to initialize a
trial and then hold the position for a short period. Visual or auditory stimuli are delivered, and
based on these stimuli, the animal must choose to poke one of the side ports. If the correct

response is chosen, the animal gets water reward from a lick tube in the response port, otherwise
a timeout punishment is applied. This training process is controlled by Bpod, an Arduino
microcontroller that interfaces with the three ports. Data from the response ports as well as video
recordings from an overhead camera are acquired simultaneously as the animal is trained.

The entire apparatus is orchestrated by a master program that coordinates the RFID sorting
device, the Bpod system, synchronized video recording, data management and logging
(**Supplementary Fig. 1e**). The program monitors the amount of water each animal consumes per
day and regulates the time each animal can spend in the training box per session. In addition, the
software updates the training protocol for each animal based on its performance, for example
switching to a harder task once a simpler one has been mastered (**Supplementary Fig. 2**). This
lets each animal learn at its own pace.

The apparatus can be assembled at a materials cost of $1500-2500, with the cheaper option using
a Raspberry Pi computer as the controller (**Supplementary Fig. 1f** and **Supplementary Table
1**). Compared with designs in which each animal is automatically trained in its own home
cage[15,17], the system saves considerable space. Because housing and training are independent
modules, the same system can be used for diverse training environments.

We tested the automated RFID sorting and animal training system by training group-house mice
to learn a variety of decision-making tasks, following similar procedures as reported
previously[11,12] (**Supplementary Fig. 2** and Online methods). The training period lasted 28 days,
with up to five mice in the common home cage. Each animal occupied the training box for 3-4
hours per day (13-15% of the 24 hours) throughout the entire training period (**Figs. 2a**, **2b** and
Supplementary Fig. 3). For a sample cohort of four animals trained in sessions of 90 trials each,
we found that the behavior box was occupied most of the time, with brief empty intervals of <10
min (**Figs. 2c, 2d** and **2e**). Each animal was trained for over 900 trials (10 sessions), and
consumed more than 1.9 mL of water per day (**Fig. 2f**). Interestingly there was no circadian
pattern to the animals' training activity, even though the setup was illuminated on a daily light
cycle (12 h on / 12 h off) (**Fig. 2g**). As observed previously, it appears that animals working for a
goal can avoid circadian modulation of the locomotor pattern[24,25].

A generalized linear model accurately predicts decision-making during training
In a decision-making task, an animal is asked to associate distinct stimuli with distinct responses.
Although this is the ultimate goal, during learning, it is often observed that the animal begins by
basing its decisions on unrelated input variables and gradually switches to using the stimulus
variables that actually predict reward. We define a policy as a mapping of these variables to the
animal's decisions. A fundamental goal in the study of learning is to infer what policy the animal
follows at any given time and to determine how the policy evolves with experience.

We applied a generalized linear model (GLM) to map factors relevant to the animal's decision-
making to its choices through logistic regression. A common way to build such a GLM is by
fitting data of an entire session[16,26]. However, this loses resolution in single trials within the
session. During learning, a change of policy can happen at each trial. Thus, we developed the
model to make trial-by-trial choice predictions based on various factors the animal might
plausibly use. The model works in an iterative two-step process (**Fig. 1b**). In the prediction step,
the model makes a prediction for the next decision based on the input factors. Once the outcome
of the animal's decision is observed, an error term between the model's prediction and the

observation is computed. This error, after weighting by a reward factor and a temporal discount
factor, is fed back to the loss function. In the update step, the model is updated by minimizing
the regularized loss function. This iteration happens after every trial. The temporal discount
factor accounts for the possibility that the most recent trials impact the current decision more
than remote trials. The reward factor accounts for the fact that water rewards and timeout
punishments may have effects of different magnitude on the updates of the animal's policy.
151
We illustrate the utility of this model by fitting results from an easy visual task, in which one of
the two choice ports lights up to indicate the location of the reward, and the optimal policy is to
simply poke the port with the light (**Supplementary Fig. 2a, 2a'** and **2a''**). All the mice
eventually reached a >83% performance level, comparable to what mice achieve in similar
tasks[19,27]. The GLM makes a prediction for the outcome of each trial based on a weighted
combination of several input variables: the current visual stimulus, a constant bias term, and
three terms representing the history of previous trials (**Fig. 3a**). These inputs from a previous
trial include the port choice, whether that choice was rewarded, and a term indicating the
multiplicative interaction between the choice and reward (Choice x Reward). This term supports
a strategy called win-stay-lose-switch (WSLS), which chooses the same port if it was rewarded
previously and the opposite one if not. Since a GLM cannot multiply two inputs, we provided
this interaction term explicitly. Each of the above terms has a weight coefficient that can be
positive or negative. For instance, a positive weight for the visual stimulus supports turning
towards the light, and a negative weight away from the light.
166
To determine the extent of trial history that affects the animal's behavior, we fitted the model to
the response data including history-dependent terms up to three previous trials. We found that
only the immediately preceding trial had an appreciable effect on the prediction accuracy, and
thus restricted further analysis to those inputs (**Fig. 3b**). The model also has three
hyperparameters (the temporal discount factor α, the reward factor r, and the regularization
factor λ), and we optimized them for each animal by grid search. We found that each animal
had a different set of hyperparameters, reflecting differences in the learning process across
individuals (**Fig. 3c**). Among the four sample mice, Animal 2 had the lowest temporal discount
factor, suggesting that it weighed recent trials more heavily and updated the policy more quickly.
Indeed, this is the animal that learned the fastest among the four (**Fig. 3d**).
177
Predictions from the iterative GLM matched ~80% of the animals' actual choices (**Fig. 3f**), and
the predicted accuracy of each animal captured the actual fluctuations of its learning curve (**Fig.
3e** and **Supplementary Fig. 4**). We compared the performance of the GLM with two other
modeling approaches (Online methods). The first model was fit to the animal's average
performance in the task; its trial-by-trial match of the animal's actual choices was only ~59%
(**Fig. 3g**). The second model was a logistic regression fitted to data in a sliding window of N
trials. This sliding window model performed worse than the iterative GLM when the window
size was small ($N = 20$ and 30 trials, **Fig. 3h**); for larger windows the performance was
comparable. Overall, the iterative model is advantageous because it makes predictions online as
every trial occurs and adapts dynamically to the growing data set.
188
Individual learning policies can be inferred from iterative GLM fitting
The iterative GLM serves to infer what policy the animal follows in making decisions. The linear
weight of each input term reflects its relative importance for the decision. By following this
weight vector across trials one obtains a policy matrix that documents how the animal's policy

changes during learning (**Figs. 1b** and **4c**). To test that the model can correctly capture a time-
varying policy, we simulated decision-making data from a ground truth policy that changed at a
certain frequency, including a certain level of noise in the behavioral output (**Fig. 4a**). Over a
wide range of policy change frequencies and noise levels, the GLM was able to capture the
ground truth policy (**Figs. 4a** and **4b**). In addition, different values of policy change frequency
and noise levels led to different sets of hyperparameters fitted from the model, showing that the
GLM can adapt to individuals with diverse learning characteristics (**Supplementary Figs. 5a-e**).

We then recovered the policy matrix of each animal from the GLM fits. All four animals started
with the non-optimal policy of WSLS. Subsequently each animal followed its own learning
process (**Fig. 4c**): Animal 2 had a clear bias towards the right port at the beginning but it rapidly
found the optimal policy of following the light. The other three animals were slower learners.
Animal 3 and Animal 4 followed similar processes to converge to the optimal policy. Animal 1
was distinct from the others. At the early stages, it had a strong bias towards the left port and it
made decisions based on whether the previous choice was rewarded.

We further validated the transition between policies during learning by analyzing the first and
last sessions of each animal and counting how many choices could be explained by each policy
(**Fig. 4d**). Indeed, we found a clear switch from the (non-optimal) WSLS policy to the (optimal)
stimulus-based policy (**Fig. 4e** and **Supplementary Fig. 5f**). The animals might have been
biased towards the WSLS strategy by a shaping method we used during training, which offered
the animal a repeat of the same stimulus every time it made a mistake (Online methods). To test
whether these correlations in the trial sequence influenced the final policy we performed two
additional analyses. First, we only included trials following a correct trial, and performed logistic
regression on these trials for each session. This analysis showed that at least on these trials, all
the animals based their decisions on the light stimulus by the end of learning (**Supplementary
Fig. 6a**). Second, we compared the error rate on trials following an incorrect choice with that
following a correct one. We found no significant difference between the two error rates during
the last session (**Supplementary Figs. 6b** and **6c**), suggesting that the animals treated these two
types of trials identically.

Automated movement tracking reveals fine structure of behavioral responses
Thus far the report has focused on the animal's responses only as sensed by the nose pokes into
response ports. The GLM fits of those responses already revealed differences in policy across
individuals. To gain further insight into these individual preferences, it is essential to track each
animal's behavior along the way from stimuli to responses[10]. We thus employed computer vision
software to automatically, quantitatively and accurately assess each animal's behavior during
decision-making. Having compared several tracking algorithms, we eventually used
DeepLabCut[45], a deep-learning based program, because it is easy to use and accurate in
identifying body landmarks (**Figs. 1c** and **5a**). We identified two body landmarks: the nose and
the centroid of the animal, and further calculated the orientation as the angle of the line
connecting the centroid and the nose (**Fig. 5a, Supplementary Video 5**).

To illustrate use of these behavioral trajectories, we focus on the period of the visual choice task
where the animal reports its decision: from the time it leaves the center port to when it pokes one
of the side ports. The trials fall into four groups based on location of the stimulus and the
response. As expected, the trajectories of position and orientation clearly distinguish left from
right choices (**Figs. 5b** and **5d**). Interestingly, the trajectories also reveal whether the decision

was correct: On incorrect decision, the trajectories reversed direction after ~0.5 s, because the animal quickly turned back to the center after finding no reward in the chosen port (**Figs. 5c, 5e** and **Supplementary Video 5**). A linear kernel support vector machine (SVM), trained to predict the category of each trial from a 1 s trajectory, was able to correctly distinguish correct and incorrect choices with an accuracy of over 90% (**Supplementary Fig. 7**). In addition, many of the trajectories were highly asymmetric and again revealed differences across individuals. For instance, Animal 2 and Animal 4 started from a location close to the right port, Animal 1 closer to the left port (**Fig. 5c**). This asymmetry correlates with the bias revealed by the iterative GLM: each animal prefers to select the port closer to its body location.

Unsupervised behavioral analysis reveals moments of hesitation
Whereas the supervised learning discussed above relies on prior classification of stimuli and responses, an unsupervised analysis has the potential to discover unexpected structures in the animal's behavior[28]. We thus performed an unsupervised classification of the behavioral trajectories.

After subjecting all the trajectories of a given animal to principal component analysis (PCA) we projected the data onto the top three components, which explained over 95% of the variance (**Figs. 6a** and **6b**). Importantly, without any labels from trial types, these three PCs captured meaningful features that differentiated the animal's responses. The first PC separated movements to the left from those to the right. The third PC captured the turning-back behavior after an incorrect choice. The second PC captured different baseline positions. Each animal has its own preference for a baseline position somewhere off the midline of the chamber (**Supplementary Fig. 8a**).

We also projected the trajectories into 2 dimensions using a non-linear embedding method, t-distributed stochastic neighbor embedding[29,30] (t-SNE). Unlike PCA, this graph prioritizes the preservation of local structures within the data instead of the global structure[30]. In the t-SNE space the trajectories formed clear clusters (**Fig. 6c**). Most of the clusters are dominated by one of the decision categories (**Fig. 6c** and **Supplementary Fig. 8b**). Interestingly, we found clusters in Animals 2, 3, and 4, in which the centroid trajectories were flat, unlike the trajectories of the four decision categories (**Fig. 6d**), suggesting that animals hesitated in these trials and made decisions only after a delay. Indeed, in trials flagged by these clusters, the animals had longer reaction times (**Fig. 6e**). Furthermore, such hesitating responses were more common following an incorrect trial (**Fig. 6f**); they may reflect a behavioral adjustment to prevent further mistakes[31].

DISCUSSION
Despite the fact that rodents can be trained to perform interesting decision-making tasks[7-10], the learning progress of individual animals has rarely been addressed. Doing so requires training and observing many animals in parallel under identical conditions, and the ability to analyze the decision policy of each animal on a trial-by-trial basis. To meet these demands, we developed Mouse Academy, an integrated platform for automated training and behavior analysis of individual animals.

We demonstrate here that Mouse Academy can train group-housed mice in an automated and highly efficient manner while simultaneously acquiring decision-making sequences and video recordings. Automated animal training has been of great interest in recent years and efforts have focused on two directions. In one design, multiple animals are trained in parallel within stacks of

training boxes. This requires a technician to transfer animals from their home cages to the
behavior boxes[14-16]. Such animal handling has been reported to introduce additional
variability[20,21], and even the mere presence of an experimenter can influence behavioral
outcomes[32]. Thus, eliminating the requirement for human intervention, as in Mouse Academy,
likely reduces experimental variation. In another design, a training setup is incorporated within
the animals' home cage[17-19]. By contrast, Mouse Academy separates the functions of housing and
training, and that modular design allows easy adaptation to a different purpose. For instance, one
can replace the 3-port discrimination box with a maze to study spatial navigation learning[33,34], or
with an apparatus for training under voluntary head-fixation[35]. In each case, a single training
apparatus can serve many mice, potentially from multiple home cages.
299
To understand how an animal's decision-making policies change in the course of learning, we
developed a trial-by-trial iterative GLM. The evolution of the model is similar to online machine
learning[36] in which the data are streamed in sequentially, rather than in batch mode. The linear
nature of the model supports a straightforward definition of the animal's decision policy, namely
as the vector of weights associated with different input variables. In addition, the simple linear
structure allows rapid execution of the algorithm, which favors its use in real-time closed-loop
behavior experiments. The model also allows several parametric adjustments. One specifies how
much the recent trials are weighted over more distant ones in shaping the animal's policy.
Another rates the relative influence of reward versus punishments. Fitting these parameters to
each animal already revealed differences in learning style. This model can have a broader use
beyond mouse decision-making, for instance to track the progress of human learners from their
answers to a series of quizzes[37].
312
Finally we analyzed behavioral trajectories of individual animals. The behavioral trajectories can
reveal intricate aspects of the animal's decision process that are hidden from a mere record of the
binary choices. The large data volume again calls for automated analysis, and both supervised
machine learning methods[28,38,39] and unsupervised classification[28-30,40] have been employed for
this purpose. Unsupervised analysis is not constrained by class labels, and can identify hidden
structure in the data in an unbiased manner. In the present case, we discovered a motif wherein
the animal hesitates on certain trials before taking action.
320
321   Mouse Academy can be combined with chronic wireless recording[41,42], to allow synchronized
322   data acquisition of neural responses. Researchers can seek correlations between neural activity
323   and the policy matrix or even the behavioral trajectories. This will open the door to a mechanistic
324   understanding of how neural representations and dynamics change in the course of animal
325   learning.
326
327   **ONLINE METHODS**
328   **Animals**
329   Subjects were C57BL/6J male mice aged 8-12 weeks. All experiments were conducted in
330   accordance with protocols approved by the Institutional Animal Care and Use Committee of the
331   California Institute of Technology.
332
333   **Hardware setup**
334   The hardware setup comprises a behavioral training box, an engineered home cage, and a radio
335   frequency identification (RFID) sorting system, which allows animals to move between the
336   home cage and the training box. These components are coordinated by customized software.

337
The design file for the behavior box was modified from that of Sanworks LLC
(https://github.com/sanworks/Bpod-CAD) using Solid-Works computer-aided design software
and the customized behavioral training box was manufactured in the lab. The behavior box is
controlled by a Bpod state machine (r0.8, Sanworks LLC). To monitor the animal's behavior, an
IR webcam (Ailipu Technology or OpenMV Camera M7) is installed above the behavior box.
The behavior box and the webcam are placed within a light- and sound-proof chamber. The
chamber is made of particle board with walls covered by acoustic foam. A tunnel made of red
plastic tubes connects the behavior box to a home cage (**Supplementary Fig. 1b)**.
346
347 For the RFID access control system, an Arduino Mega 2560 microcontroller is connected with
348 three RFID readers (ID-12LA, Sparkfun) with custom antenna coils spaced along the access
349 tunnel. The microcontroller controls two generic servo motors fitted with plastic gates to grant
350 individual access to the training box (**Supplementary Fig. 1a)**.
351
The microcontroller identifies each animal by its implanted RFID chip and permits only one
animal to go through the tunnel connecting the home cage and the behavioral training box
(**Supplementary Fig. 1c)**. It also communicates the animal's identity to a master program
running on a PC or Raspberry Pi (Matlab or Python). The master program coordinates the
following programs: Bpod (https://github.com/sanworks/Bpod), synchronized video recording,
data management and logging. A repository containing the design files, the firmware code for the
microcontroller, and the software can be found in
https://github.com/muqiao0626/Mouse_Academy.
360
Behavior training
The training procedures of mice to perform a selective attention task are similar to those
previously reported[11,12]. Mice were water restricted for seven days before training, and
habituated in the automated training system to collect reward freely for several sessions. Then
the mice were trained in sessions, each of which was made of 90 trials, to collect water rewards
by performing two alternative forced choice tasks. Briefly, the animal had to nose-poke one of
two choice ports based on the presented stimuli. If the decision was correct, 10% sucrose-
sweetened water (3 µL) was delivered to the animal. For incorrect responses, the animal was
punished with a five-second timeout. Following an incorrect response, the animal was presented
with the identical trial again; this simple shaping procedure helps counter-act biases in the
behavior.
Over 28 days of training the animals learned increasingly complex tasks, from visual
discrimination to a two-modality cued attention switching task[11,12]. The training progressed
through six stages (**Supplementary Fig. 2**):
1. A simple visual task: In this task, the animal initiates a trial by poking the center port and
holding the position for 100 ms. Then either the left or right side port light up briefly until
the animal moves away from the center port. The animal must then poke one of the two
side ports within the decision period of 10 s. Choice of the port flagged by the light leads
to a water reward, and choice of the other port leads to a time-out period during which no
trials can be initiated. Data presented in the main text are from this stage of training only.
2. A simple auditory task: As Stage 1, except that the stimulus was white noise sound either
the left or the right side to flag the reward port.
3. A cued single-modality (visual or auditory) switching task: Blocks of 15 trials consisting
of single-modality (visual or auditory) stimulus presentation. Each block was like stages

1 or 2, except that the trial type was indicated by a 7 kHz (visual) or 18 kHz (auditory)
pure tone.
4. A cued single- and double-modality switching task: Like stage 3, but distracting trials
were introduced in which both visual and auditory stimuli were present, but only one of
the modalities was relevant to the decision. The relevant modality was again indicated by
the pure tone cues. In repeating blocks, four types of trials were presented: a. five visual-
only trials; b. ten 'attend to vision' trials with auditory distractors; c. five auditory-only
trials; d. ten 'attend to audition' trials with visual distractors. During the training, the time
that the animal had to hold in the center port was gradually increased to 0.5 s, and the
duration of the stimuli was gradually shortened to 0.2 s.
5. A cued double-modality switching task: Like stage 4 except that the single-modality trials
were removed, and the block length was gradually shortened to three trials.
6. A selective attention task: Like stage 5, but the block structure was abandoned and all
eight possible trial types were randomized: (audition vs vision) x (sound left or right) x
(light left or right).
400
401 **Iterative generalized linear model**
402 We modeled the animal's choice probability by a logistic regression. At each trial number t, the
403 choice probability is defined as

$$p\left(y_t = 1 \middle| \mathbf{w}_{t-1}\right) = \frac{1}{1 + \exp\left(-\mathbf{w}_{t-1}^T \mathbf{x}_t\right)}$$

404   (1)

$$p\left(y_t = -1 \middle| \mathbf{w}_{t-1}\right) = 1 - p\left(y_t = 1 \middle| \mathbf{w}_{t-1}\right)$$

405
406 where y_t indicates the binary choice of the animal (1 = right, –1 = left), $\mathbf{x}_t$ is the vector of input
407 factors on trial t, and $\mathbf{w}_{t-1}$ is the vector of weights for these factors obtained from fitting up to
408 the preceding trial.

409 The prediction y_t^* for the animal's choice is simply that with the higher model probability:

$$y_t^* = \underset{y \in \{-1,+1\}}{\arg\max} \, p\left(y \middle| \mathbf{x}_t, \mathbf{w}_{t-1}\right) \tag{2}$$

411 After observing the animal's actual choice z_t, the cross-entropy error E_t between the
412 observation and model prediction is calculated as

$$E_t = -\log p\left(z_t \middle| \mathbf{x}_t, \mathbf{w}_{t-1}\right) \tag{3}$$

414 We weight the error term by a reward factor R_t, and apply exponential temporal smoothing to
415 get the loss function L_t:

$$L_t = R_t E_t + \alpha L_{t-1} \tag{4}$$

417 where α is the smoothing discount factor accounting for the effect that distant trials have less
418 impact on decision-making than immediately preceding trials, and R_t is defined as

$$R_t = \begin{cases} 1, \text{if the choice is rewarded} \\ r, \text{otherwise} \end{cases} \tag{5}$$

420 The values of R_t for rewarded and unrewarded trials may be different, accounting for the fact
421 that rewards and punishments may have different effects on learning. For each time point, the
422 weights in the model are determined by minimizing the loss function subject to L1 (lasso)
423 regularization, namely

$$\mathbf{w}_t = \arg\min_{\mathbf{w}} \left(L_t + \lambda \|\mathbf{w}\|_1 \right) \qquad (6)$$

425 Then $\mathbf{w}_t$ is used for prediction of the next trial. For subsequent analysis, we only used
426 predictions starting at the 15th trial. The three hyperparameters for the temporal discount factor
427 α, the reward factor r, and the regularization factor λ were selected by grid search.
428
429 To fit the decision-making sequences of the simple visual task, we included the following terms
430 in the input vector $\mathbf{x}_t$:
     1. Visual_stimulus: +1 = light on right, –1 = light on left.
     2. Bias: A constant value of +1. The associated weight determines whether the animal
        favors the left (negative) or the right (positive) port.
     3. Choice_back_n: The choice the animal made n trials ago (+1 = right, –1 = left).
     4. Reward_back_n: The reward the animal received n trials ago (+1 = reward, –1 =
        punishment).
     5. Choice x Reward_back_n: The product of terms 4 and 5. This term corresponds to the
        win-stay-lose-switch (WSLS) strategy of repeating the last choice if it was rewarded and
        switching if it was punished.

441 To determine the extent of history-dependence of the animal's decisions, we fitted the model
442 including terms 3-5 from up to three previous trials (n = 1, 2, 3), and found that only the
443 immediately preceding trial had an appreciable effect on the model's prediction accuracy. For the
444 subsequent analysis, we therefore included terms 3-5 for the preceding trial (n = 1).
445
446 We compared the iterative generalized linear model (GLM) with two other models. The first
447 only captures the animal's average performance over all trials. If the fraction of the correct
448 responses is z, then the model simply predicts a correct response with probability z, and an error
449 with probability $1-z$. Thus, the fraction of trials where the prediction matches the observation is
450 $z^2 + (1-z)^2$.
451
452 The second model is a sliding window logistic regression. To make a prediction for trial t, we
453 fitted the logistic model presented above (Eqns 1-2) to the preceding n trials. The loss function is

$$L_t = -\sum_{i=t-n}^{t-1} \log p\left(z_t \mid \mathbf{x}_t, \mathbf{w}_{t-1}\right) \qquad (7)$$

455 and the weights are again optimized as in Eqn 6.
456
Recovering policy matrices from simulated data
To test the model's ability in recovering policy matrices, we trained the model on data generated
from pre-defined ground truth policies. The ground truth policies changed every 10 trials, 30
trials, or 90 trials. Binary choices were simulated with different noise levels using the algorithm
'epsilon-greedy': with a probability of epsilon, the simulator made a random choice and with a

probability of 1-epsilon it chose the action indicated by the ground truth policy. The noise levels
(epsilon values) ranged from 0 to 0.6. The similarity between the recovered policy and the
ground truth policy was evaluated by the cosine between the recovered weight vector and the
ground truth weight vector.
466
Supervised and unsupervised analysis of behavioral trajectories
We annotated two body landmarks, the nose and the centroid, on ~100 frames of the video, and
used them to train DeepLabCut[45]. Tested on a separate set of annotated frames, more than 85%
the nose positions are inferred within an error radius of 0.25 cm, and more than 85% of the
centroid positions are inferred within an error radius of 0.4 cm. From the nose and the centroid,
we calculated the orientation as the angle of the line connecting the centroid and the nose. For
each trial, the centroid and orientation were extracted for n frames ($n = 30$ (1 s) in most cases),
thus the data dimension for each trial is $3n$ (the two centroid coordinates and the orientation).
475
To determine whether the behavioral trajectories contain information about the decision
categories, a support vector machine (SVM) with a linear kernel was trained for each decision
category. The training set was labelled with the decision category based on information about the
visual stimulus and the animal's choice (for example, "Stim: R, Choice: L" means that the light
is on the right and the animal chooses the left port). Performance of the trained SVM was
examined by prediction accuracy on the test set, and the F1 score, which is the harmonic mean of
precision and recall:

$$F_1 = 2 \cdot \frac{precision \cdot recall}{precision + recall} = \frac{2 \cdot true\_positive}{2 \cdot true\_positive + false\_positive + false\_negative} \quad (8)$$

The performance was computed as the average across 10 repeated analyses (**Supplementary**
Fig. 7).
486
We performed a non-linear embedding method, t-distributed stochastic neighbor embedding (t-
SNE) analysis as previously described[29,30]. Briefly, the trajectory data of each trial were
projected into a 2D t-SNE space. Point clouds on the t-SNE map represented candidate clusters.
Density clustering identified these regions. We then plotted trajectories and reaction time
distributions to confirm that the clusters were distinct from each other. A repository of the
analysis scripts can be found in https://github.com/tonyzhang25/MouseAcademyBehavior.
493

AUTHOR CONTRIBUTIONS
M.Q. and M.M. designed the study; M.Q. and S.S. constructed the hardware setup and wrote the
controlling software; M.Q. performed experiments and collected data for analysis; M.Q.
developed the iterative generalized linear model with input from P.P. and M.M; C.S.
implemented animal tracking and provided help in analyzing the mouse trajectories; T.Z.
analyzed behavioral trajectories with input from M.Q., P.P. and M.M; M.Q. and M.M. wrote the
manuscript with comments from all authors.

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COMPETING FINANCIAL INTERESTS
The authors declare no competing financial interests.
511
512 **DATA AVAILABILITY**
513 The datasets analyzed during the current study are available in
514 https://drive.google.com/open?id=1gkPbqGYKPGs7Rx1WNmubQW0dKyYE5YVR
515

REFERENCES
1   Spear, N. E., Miller, J. S. & Jagielo, J. A. Animal memory and learning. *Annu Rev*
Psychol **41**, 169-211, doi:10.1146/annurev.ps.41.020190.001125 (1990).
2   Branson, K., Robie, A. A., Bender, J., Perona, P. & Dickinson, M. H. High-throughput
ethomics in large groups of Drosophila. *Nature methods* **6**, 451-457,
doi:10.1038/nmeth.1328 (2009).
3   Honegger, K. & de Bivort, B. Stochasticity, individuality and behavior. *Current biology :*
CB **28**, R8-R12, doi:10.1016/j.cub.2017.11.058 (2018).
4   Matzel, L. D. & Sauce, B. Individual differences: Case studies of rodent and primate
intelligence. *J Exp Psychol Anim Learn Cogn* **43**, 325-340, doi:10.1037/xan0000152
(2017).
5   Atkinson, J. W. & Raphelson, A. C. Individual differences in motivation and behavior in
particular situations. *J Pers* **24**, 349-363 (1956).
6   Stern, E. Individual differences in the learning potential of human beings. *Npj Science of*
Learning **2** (2017).
7   Luo, L., Callaway, E. M. & Svoboda, K. Genetic Dissection of Neural Circuits: A
Decade of Progress. *Neuron* **98**, 865, doi:10.1016/j.neuron.2018.05.004 (2018).
8   Carandini, M. & Churchland, A. K. Probing perceptual decisions in rodents. *Nature*
neuroscience **16**, 824-831, doi:10.1038/nn.3410 (2013).
9   Gomez-Marin, A., Paton, J. J., Kampff, A. R., Costa, R. M. & Mainen, Z. F. Big
behavioral data: psychology, ethology and the foundations of neuroscience. *Nature*
neuroscience **17**, 1455-1462, doi:10.1038/nn.3812 (2014).
10   Juavinett, A. L., Erlich, J. C. & Churchland, A. K. Decision-making behaviors: weighing
ethology, complexity, and sensorimotor compatibility. *Current opinion in neurobiology*
49, 42-50, doi:10.1016/j.conb.2017.11.001 (2018).
11   Schmitt, L. I. *et al.* Thalamic amplification of cortical connectivity sustains attentional
control. *Nature* **545**, 219-223, doi:10.1038/nature22073 (2017).
12   Wimmer, R. D. *et al.* Thalamic control of sensory selection in divided attention. *Nature*
526, 705-709, doi:10.1038/nature15398 (2015).
13   Akrami, A., Kopec, C. D., Diamond, M. E. & Brody, C. D. Posterior parietal cortex
represents sensory history and mediates its effects on behaviour. *Nature* **554**, 368-372,
doi:10.1038/nature25510 (2018).
14   Uchida, N. & Mainen, Z. F. Speed and accuracy of olfactory discrimination in the rat.
Nature neuroscience **6**, 1224-1229, doi:10.1038/nn1142 (2003).
15   Scott, B. B., Brody, C. D. & Tank, D. W. Cellular resolution functional imaging in
behaving rats using voluntary head restraint. *Neuron* **80**, 371-384,
doi:10.1016/j.neuron.2013.08.002 (2013).
16   Busse, L. *et al.* The detection of visual contrast in the behaving mouse. *The Journal of*
neuroscience : the official journal of the Society for Neuroscience **31**, 11351-11361,
doi:10.1523/JNEUROSCI.6689-10.2011 (2011).

17   Poddar, R., Kawai, R. & Olveczky, B. P. A Fully Automated High-Throughput Training
System for Rodents. *PloS one* **8**, doi:ARTN e83171
10.1371/journal.pone.0083171 (2013).
18   Silasi, G. *et al.* Individualized tracking of self-directed motor learning in group-housed
mice performing a skilled lever positioning task in the home cage. *Journal of*
Neurophysiology **119**, 337-346, doi:10.1152/jn.00115.2017 (2018).
19   Erskine, A. B., T.; Herb, J. T.; Schaefer, A. AutonoMouse: High throughput automated
operant conditioning shows progressive behavioural impairment with graded olfactory
bulb lesions. https://www.biorxiv.org/content/early/2018/03/29/291815 (2018).
20   Crabbe, J. C., Wahlsten, D. & Dudek, B. C. Genetics of mouse behavior: interactions
with laboratory environment. *Science* **284**, 1670-1672 (1999).
21   Hurst, J. L. & West, R. S. Taming anxiety in laboratory mice. *Nature methods* **7**, 825-
826, doi:10.1038/nmeth.1500 (2010).
22   Daw, N. D. in *Decision Making, Affect, and Learning: Attention and Performance XXIII*
(Oxford University Press, 2011).
23   Winter, Y. & Schaefers, A. T. A sorting system with automated gates permits individual
operant experiments with mice from a social home cage. *J Neurosci Methods* **196**, 276-
280, doi:10.1016/j.jneumeth.2011.01.017 (2011).
24   Daan, S. *et al.* Lab mice in the field: unorthodox daily activity and effects of a
dysfunctional circadian clock allele. *J Biol Rhythms* **26**, 118-129,
doi:10.1177/0748730410397645 (2011).
25   Hut, R. A., Pilorz, V., Boerema, A. S., Strijkstra, A. M. & Daan, S. Working for food
shifts nocturnal mouse activity into the day. *PloS one* **6**, e17527,
doi:10.1371/journal.pone.0017527 (2011).
26   Seidemann, E. Neuronal mechanisms mediating conversion of visual signals into
perceptual decisions in a direction discrimination task. *Ph.D. dissertation, Stanford*
University. (1998).
27   Burgess, C. P. *et al.* High-Yield Methods for Accurate Two-Alternative Visual
Psychophysics in Head-Fixed Mice. *Cell Rep* **20**, 2513-2524,
doi:10.1016/j.celrep.2017.08.047 (2017).
28   Gris, K. V., Coutu, J. P. & Gris, D. Supervised and Unsupervised Learning Technology
in the Study of Rodent Behavior. *Front Behav Neurosci* **11**, 141,
doi:10.3389/fnbeh.2017.00141 (2017).
29   Berman, G. J., Choi, D. M., Bialek, W. & Shaevitz, J. W. Mapping the stereotyped
behaviour of freely moving fruit flies. *J R Soc Interface* **11**, doi:10.1098/rsif.2014.0672
(2014).
30   Todd, J. G., Kain, J. S. & de Bivort, B. L. Systematic exploration of unsupervised
methods for mapping behavior. *Phys Biol* **14**, 015002, doi:10.1088/1478-
3975/14/1/015002 (2017).
31   Botvinick, M. M., Braver, T. S., Barch, D. M., Carter, C. S. & Cohen, J. D. Conflict
monitoring and cognitive control. *Psychol Rev* **108**, 624-652 (2001).
32   Sorge, R. E. *et al.* Olfactory exposure to males, including men, causes stress and related
analgesia in rodents. *Nature methods* **11**, 629-632, doi:10.1038/nmeth.2935 (2014).
33   Barnes, C. A. Memory deficits associated with senescence: a neurophysiological and
behavioral study in the rat. *J Comp Physiol Psychol* **93**, 74-104 (1979).
34   Olton, D. S., Walker, J. A. & Gage, F. H. Hippocampal connections and spatial
discrimination. *Brain Res* **139**, 295-308 (1978).

35 Aoki, R., Tsubota, T., Goya, Y. & Benucci, A. An automated platform for high-
throughput mouse behavior and physiology with voluntary head-fixation. *Nat Commun* **8**,
1196, doi:10.1038/s41467-017-01371-0 (2017).
36 Shalev-Shwartz, S. Online Learning and Online Convex Optimization. *Foundations and*
Trends in Machine Learning **4**, 107-194 (2012).
37 Piech, C. S., J.; Huang, J.; Ganguli, S.; Sahami, M.; Guibas, L.; Sohl-Dickstein, J. Deep
Knowledge Tracing. https://arxiv.org/abs/1506.05908 (2015).
38 Hong, W. *et al.* Automated measurement of mouse social behaviors using depth sensing,
video tracking, and machine learning. *Proc Natl Acad Sci U S A* **112**, E5351-5360,
doi:10.1073/pnas.1515982112 (2015).
39 Egnor, S. E. & Branson, K. Computational Analysis of Behavior. *Annu Rev Neurosci* **39**,
217-236, doi:10.1146/annurev-neuro-070815-013845 (2016).
40 Wiltschko, A. B. *et al.* Mapping Sub-Second Structure in Mouse Behavior. *Neuron* **88**,
1121-1135, doi:10.1016/j.neuron.2015.11.031 (2015).
41 Szuts, T. A. *et al.* A wireless multi-channel neural amplifier for freely moving animals.
Nature neuroscience **14**, 263-269, doi:10.1038/nn.2730 (2011).
42 Ziv, Y. *et al.* Long-term dynamics of CA1 hippocampal place codes. *Nature*
neuroscience **16**, 264-266, doi:10.1038/nn.3329 (2013).
43 Dankert, H., Wang, L., Hoopfer, E. D., Anderson, D. J. & Perona, P. Automated
monitoring and analysis of social behavior in Drosophila. *Nature methods* **6**, 297-303,
doi:10.1038/nmeth.1310 (2009).
44 Eyjolfsdottir, E. B., K.; Yue, Y.; Perona, P. Learning recurrent representations for
hierarchical behavior modeling. https://arxiv.org/abs/1611.00094 (2016).
45 Mathis, A. *et al.* DeepLabCut: markerless pose estimation of user-defined body parts
with deep learning. *Nature neuroscience* **21**, 1281-1289, doi:10.1038/s41593-018-0209-y
(2018).
629

Figure 1

a

IR camera

RFID antennas

Behavior training
box with three ports

Mouse implanted
with a RFID chip

Home cage

Motorized gates

RFID-tagged mice group housed in the home cage

RFID based animal sorting

Animal training and behavior profiling

b

At each trial

Relevant factors x

w

Update

Prediction y

Loss function L   α

Iterative GLM

Observed choice and
whether it is rewarded

Weight vector
over time

Policy matrix

Trial:	1	2	3	...	N
Factor 1	0.8	0.8	0.8		1
Factor 2	0.2	0	0		0
Factor 3	0.3	0.25	0.2		0
...	...	...	...	...	...
Factor n	0.5	0.5	0.25		0

c

Video data

Deep convolutional neural nets

Features: centroid
and orientation

Animal's behavioral
trajectories

...

Correlate with
decision-making

Feature

Time

Fig. 1: Components of Mouse Academy
(a) An automated RFID sorting and animal training system. Mice implanted with RFID chips are group-housed in the home cage. The RFID sorting system identifies each mouse by its implanted chip. One animal at a time gains access to a behavioral training box. As the animal learns a task, its decision sequences and video recordings are acquired. **(b)** An iterative generalized linear model. For each trial, the model predicts the animal's choice based on the relevant factors and then evaluates the difference from the actual choice. This difference, after temporal weighting, is fed back to the loss function, which gets minimized by updating the weights of the input factors. The model produces a policy matrix in which the rows indicate the weights of the relevant factors and the columns are the trials. **(c)** An automated behavior assessment program using deep convolutional neural networks to extract the location and pose information of an animal.

Figure 2

Fig. 2: Performance of the automated training system on a sample cohort
(a) Fraction of time the behavior box was occupied by each of the four animals. **(b)** Activity trace of each animal in the behavior box for the entire training period of 28 days. Shadow indicates the dark cycle from 8pm to 8am. **(c)** Distribution of time intervals during which the behavior box is occupied or empty. **(d)** Box plot of intervals between each animal's sessions (median, quartiles, and range). **(e)** Box plot of the time spent in a session for each animal. **(f)** Averaged daily water consumption of each animal. Error bars indicate standard errors. **(g)** Circadian histograms of each animal's activity in the behavior box.

Figure 3

Fig. 3: Iterative generalized linear model and its prediction accuracy
(a) Illustration of the GLM as applied to a visual discrimination task. The model's prediction is based on the output of a logistic function whose input is the weighted sum of a visual stimulus term, a bias term, and three history-dependent terms. The stimulus can be on the left or right and the choice can be rewarded (consistent with the stimulus, indicated by a green dot) or unrewarded (opposite to the stimulus, indicated by a red dot). **(b)** Selection of the history-dependent terms based on the model prediction accuracy. Error bars indicate standard errors. **(c)** Hyperparameters for each of the animals: reward factor, discount factor, and regularization factor. The optimal values are marked with a star. **(d)** The actual performance of each animal over time in the visual task. **(e)** Performance as predicted by the GLM. **(f)** Fraction of choices predicted correctly by the GLM. **(g)** Fraction of choices predicted correctly by a simple model based on the animal's average performance in the task. **(h)** Fraction of predictions matched by the iterative GLM and the sliding window logistic regression model. Error bars indicate standard errors. **, * indicate $P < 0.01, 0.05$. Random prediction would give 50% match.

Figure 4

Fig. 4: Interpretation of policies during learning

(a) Policy vectors recovered by the iterative GLM capture the ground truth policies. The policy matrix plots in each trial (horizontal) the weights associated with each of 5 factors (vertical), encoded with a color scale (see panel c). The factors are: A = Visual_stimulus, B = Choice x Reward_back_1, C = Choice_back_1, D = Reward_back_1, E = Bias. Two examples are shown of ground truth policies used to simulate data and the corresponding trial-by-trial estimates from the GLM. Blanks in the ground truth matrix indicate instances where the simulated choice is opposite to the policy. **(b)** Similarity between the recovered policy and the ground truth, measured by the cosine between the two policy vectors. Error bars indicate standard deviation. **(c)** Policy matrices recovered for the four animals show distinct individual learning processes. Dashed rectangles highlight the first and last sessions of each animal, as enlarged in **d**. **(d)** Recovered policy matrices for the first and last sessions of each animal. **(e)** Fraction of trials explained by two candidate policies (win-stay-lose-switch and following the stimuli) in the first and last sessions. Error bars indicate standard errors. ** indicates $P < 0.01$.

Figure 5

Fig. 5: Supervised analysis using features extracted by automated behavior assessment
(a) DeepLabCut extracts the centroid and the orientation as the angle between the horizontal axis
and the line connecting the centroid and the nose. **(b)** Centroid distance along the left-right axis
vs time during the movement, for animal 1. The starting position is set to zero, positive values
indicate movement to the left, negative to the right. The 4 trial types are indicated by different
colors. **(c)** Average centroid trajectory for each animal. Shaded region indicates standard error.
(d-e) Orientation vs time, displayed as in panels b-c. Positive angle points to the left, negative to
the right.

Figure 6

a

Animal1 Animal2 Animal3 Animal4

Stim:L,Choice:L
Stim:L,Choice:R
Stim:R,Choice:R
Stim:R,Choice:L

b

PC1: 0.74   PC1: 0.82   PC1: 0.76   PC1: 0.73
PC2: 0.21   PC2: 0.13   PC2: 0.17   PC2: 0.21
PC3: 0.02   PC3: 0.02   PC3: 0.03   PC3: 0.02
PC4: 0.01   PC4: 0.01   PC4: 0.02   PC4: 0.02

c

d

e

f

Fig. 6: Unsupervised analysis of the behavior trajectories
(a) Principal component projections onto PC1 and PC2 of the centroid-vs-time trajectories from Fig 5. The 4 trial types are indicated by different colors. **(b)** The centroid trajectories corresponding to the first four principal components (PCs). The variance explained by each PC is shown in the plot legend. **(c)** Clustering trials by their trajectories using t-SNE analysis. Distinct clusters are marked with different colors for use in subsequent panels. **(d)** Averaged centroid distance vs time for each cluster, plotted as in Fig 5b. **(e)** Box plot of the reaction time for each cluster. **(f)** The error rate on the preceding trial for each cluster. Error bars indicate standard errors.

1   **Supplementary Materials**
2
3   **Supplementary Table 1. Cost Analysis of Mouse Academy Hardware**
4   **PC version**

Parts	Quantity	Supplier	Cost
Bpod State Machine	1	Sanworks	$495.00
Custom behavior box (Plastic, ports with LEDs and photo-gates, valves, etc.)	1	Port breakout boards from Sanworks	$500.00
IR Webcam	1	Ailipu Technology	$47.99
Custom light- and sound-proof chamber	1	NA	$75.00
Arduino Mega 2560 R3	1	Sparkfun	$45.95
RFID Reader ID-12LA	3	Sparkfun	$29.95
Servo - Generic	2	Sparkfun	$8.95
Custom tunnel (Plastic, holders, antennas etc.)	1	NA	$150.00
Engineered home cage	1	Caltech animal facility	$76.10
PC	1	Hewlett-Packard	$1000.00
Total cost			$2428.94

5
6   **Raspberry Pi version**

Parts	Quantity	Supplier	Cost
Bpod State Machine	1	Sanworks	$495.00
Custom behavior box (Plastic, ports with LEDs and photo-gates, valves, etc.)	1	Port breakout boards from Sanworks	$500.00
OpenMV Camera M7	1	OpenMV	$65.00
Custom light- and sound-proof chamber	1	NA	$75.00
Arduino Mega 2560 R3	1	Sparkfun	$45.95
RFID Reader ID-12LA	3	Sparkfun	$29.95
Servo - Generic	2	Sparkfun	$8.95
Custom tunnel (Plastic, holders, antennas etc.)	1	NA	$150.00
Engineered home cage	1	Caltech animal facility	$76.10
Raspberry Pi 3 Model B	1	Raspberry Pi	$35.10
Total cost			$1481.05

8 **Supplementary Table 2: mean averaged precision (mAP) and recall (mAR) at different**
9 **threshold of Intersection over Union (IoU) (related to mouse detection)**

	mAP	mAR
$0.75 < IoU < 0.95$	0.81	0.82
$IoU = 0.5$	0.99	0.99
$IoU = 0.75$	0.95	0.98

Supplementary Figure 1

Supplementary Fig. 1: Technical details of the hardware design
(a-b) Side view of the setup (**a**) packed into a light- and sound-proof box (**b**). **(c)** RFID sorting process. For an animal to enter the behavior box, only when the left and the middle detectors detect the same RFID chip, the left gate is closed and the right gate is open so that the animal can access the behavior box. For an animal to return to the home cage, only when the right and the middle detectors detect the same RFID chip, the right gate is closed and the left gate is open so that the animal can go back to the home cage. In the entry and the return processes, if the left and the middle detectors detect different RFID chips, the animals have to leave the tube and the detectors get reset afterwards. **(d)** Schematic of RFID access control circuit. **(e)** Schematic of the software controlling the devices. A master program receives input from the RFID sorting device and controls four other modules including Bpod, synchronized video recording, data management, and logging. **(f)** Top view of a Raspberry Pi version of the setup.

Supplementary Figure 2

Supplementary Fig. 2: Illustration of training procedures

Training proceeds through six stages (Online methods). The design, learning curves, and animal performance of the simple visual task **(a, a', a'')**, the simple auditory task **(b, b', b'')**, the cued single-modality (visual or auditory) switching task **(c, c', c'')**, the cued single- (visual or auditory) and double-modality (attend to vision or audition) switching task **(d, d', d'')**, the cued double-modality (attend to vision or audition) switching task **(e', e'')**, and the final selective attention task **(f, f', f'')** are shown here. **a'** displays performance data as in **Fig. 3e**. Brown and gray dashed lines indicate the performance thresholds for upgrading to the next stage and downgrading to the previous stage respectively.

Supplementary Figure 3

a

b

32   **Supplementary Fig. 3: Automated training system allows efficient use of the behavior box**
33   For a sample cohort of five animals, this shows the fraction of time each animal used the
34   behavior box (**a**) and the activity trace of each animal throughout one month.

Supplementary Figure 4

Supplementary Fig. 4: Additional analysis on the iterative generalized linear model's prediction accuracy

The actual performance and the performance predicted by the model, for each of the four animals. Note that the predictions recapitulate the more prominent fluctuations in the actual learning curves. Error bars indicate standard errors.

Supplementary Figure 5

Supplementary Fig. 5: Iterative generalized linear model captures differences between
individuals and policy changes
(a-b) Hyperparameter selection for the GLMs fitted to the simulated data generated from the
ground truth policies. Different values of policy change frequency and noise level (epsilon) lead
to different landscapes of the hyperparameters. **(c-e)** Selected temporal discount factor (**c**),
reward factor (**d**) and regularization factor (**e**) for different values of policy change frequency
and noise level (epsilon). **(f)** Fraction of the trials explained by the two policies (win-stay-lose-
switch or WSLS, and following the stimuli) in the first and last sessions, for each of the four
animals.

Supplementary Figure 6

b

c

Supplementary Fig. 6: Additional analyses of policy changes during learning
(a) Policy matrices over sessions of the four animals. Here the policy matrices are recovered
from logistic regression using only the trials following a correct response. Because the reward of
the last trial is always +1, the term Reward_back_1 is the same as Bias, and the term
RewardxChoice_back_1 is equal to Choice_back_1, so we drop them to avoid redundancy. **(b-c)**
Quantification of the error rate during the last session, comparing trials following a correct
response to those following a mistake. Averaged over all four animals (**b**) and for each of the
four animals (**c**). n.s. indicates not significant.
57

Supplementary Figure 7

a

b

Supplementary Fig. 7: Performance of the support vector machine to infer trial category from mouse trajectories

(a) Prediction accuracy of the SVMs for individual animals. **(b)** F1 score of the SVM fitted for the decision categories of each animal. Shaded region denotes standard error. x axis indicates the time starting from when the animal leaves the center port to make a choice. SVMs were trained using features up to a certain time point.

Supplementary Figure 8

Supplementary Fig. 8: Additional information on the unsupervised analysis of behavior
trajectories
(a) Scatter plot of starting positions along the left-right axis against PC2 shows correlation
between the two. Starting positions are normalized to range from 0 (the leftmost position) and 1
(the rightmost position). **(b)** t-SNE plots with colors indicating different decision categories.

Supplementary Video 1:
https://drive.google.com/open?id=1Ng5s1UhlFRdV4mZ5b1Ot7EUpaEiUXN2_
71
72 **Supplementary Video 2:**
73 https://drive.google.com/open?id=15qgqM5qOd30kajT-IQkCf8flcF80U2qV
74
75 **Supplementary Video 3:**
76 https://drive.google.com/open?id=1zqja6_3bA2jO9ap0EWd5t_Z_dVOA8FmA
77
78 **Supplementary Video 4:**
79 https://drive.google.com/open?id=1wiaaBD-sfZTudDbpRM0ByB2n9bTKyaid
80
81 **Supplementary Video 5:**
82 https://drive.google.com/open?id=1TQd6N4XW2LDtLbBxkLQezoTJfacNHEZV
83