

Multi-dimensional reward evaluation in mice

Vladislav Nachev^{1,*†§}, Marion Rivalan^{1,†§}, York Winter^{1,§}

¹ Humboldt University, Berlin, Germany

***For correspondence:** vladislav.nachev@charite.de

†These authors contributed equally to this work

Present Address: [§]Dept. of Biology, Humboldt University, Philippstr. 13, 10099 Berlin, Germany

Abstract

The experiments in this study were designed to test how mice use information from two reward dimensions (probability and volume) when deciding where to forage.

[Please provide an abstract of no more than 150 words. Your abstract should explain the main contributions of your article, and should not contain any material that is not included in the main text.]

Introduction

(Levy and Glimcher 2012) (Kacelnik and Brito e Abreu 1998) (Rosenström, Wiesner, and Houston 2016) (Rivalan, Winter, and Nachev 2017) (Matell, Kim, and Hartshorne 2014) The partial preference observed in choice experiments can be explained by profitability matching (Kacelnik 1984), which states that animals proportionally allocate their effort depending on the relative pay-off of the options.

Results

In order to test how (contradicting) information from two different dimensions is integrated and weighed, we performed a series of choice experiments (1-4, in chronological order) with mice in automated group cages (Rivalan, Winter, and Nachev 2017). The cages were outfitted with four computer-controlled liquid dispensers that delivered drinking water as a reward. During each of the 18h-long drinking sessions each mouse had access to all dispensers, but received rewards at only two of them. The two rewarding dispensers differed on one or both reward dimensions, probability and volume (Rivalan, Winter, and Nachev 2017). An overview of the differences between choice options in the different experimental conditions is given in Table 1. All experiments were conducted with three different cohorts of eight mice each. Cohort 2 was housed in a different automated group cage than cohorts 1 and 3 (See Animals, Materials, and Methods for differences between cages).

Experiment 1

In the baseline conditions rewards only differed on one dimension (the relevant dimension), but not on the other dimension (the background dimension). For example, in the BVP1 condition (read as baseline for volume at probability 1), both options had the same probability of 0.2, but one option had a volume of 4 μ L and the other, a volume of 20 μ L (Table 1). Based on previous experiments (Rivalan, Winter, and Nachev 2017), we expected a baseline difference between 4 μ L and 20 μ L volumes to result in a similar discrimination performance (relative preference for the superior option) compared to a baseline difference between probabilities 0.2 and 0.5. In the C (congruent) condition one option was superior to the other on both dimensions. Finally, in the I (incongruent) condition each of the two options was superior to the other

on one of the two reward dimensions. Since the differences on both dimensions were chosen to be comparable, we expected the mean discrimination performance in the incongruent condition to be at chance level (0.5). In experiment 1 and in all subsequent experiments, each mouse had its own individual sequence of conditions, but each condition was followed by a reversal in the next drinking session, with a spatial inversion of the two rewarding dispensers. In order to investigate how the two reward dimensions contributed towards choice, we looked at the contrasts between the baselines (when only one dimension was relevant) to the conditions when the two dimensions were congruent or incongruent to each other. We used equivalence tests (Lakens 2017) with an *a priori* smallest effect size of interest (sesoi) of 0.1, i.e. we only considered absolute differences of at least 0.1 percentage points to be of biological relevance. Smaller differences, regardless of their statistical significance using other tests, were considered to be trivial.

Table 1: Overview of the experimental conditions in all four experiments.

experiment ^a	condition	option A			option B			A/B
		volume ^b	probability	return ^c	volume ^b	probability	return ^c	relative return
1	BPV1	4	0.2	0.8	4	0.5	2.0	0.40
1	BPV2	20	0.2	4.0	20	0.5	10.0	0.40
1	BVP1 ^d	4	0.2	0.8	20	0.2	4.0	0.20
1	BVP2	4	0.5	2.0	20	0.5	10.0	0.20
1	C	4	0.2	0.8	20	0.5	10.0	0.08
1	I	4	0.5	2.0	20	0.2	4.0	0.50
2	BPV1	4	0.2	0.8	4	1.0	4.0	0.20
2	BPV2	20	0.2	4.0	20	1.0	20.0	0.20
2	BVP2	4	1.0	4.0	20	1.0	20.0	0.20
2	C	4	0.2	0.8	20	1.0	20.0	0.04
2	I	4	1.0	4.0	20	0.2	4.0	1.00
3	PV1	4	0.2	0.8	4	0.5	2.0	0.40
3	PV2	10	0.2	2.0	10	0.5	5.0	0.40
3	PV3	15	0.2	3.0	15	0.5	7.5	0.40
3	PV4	20	0.2	4.0	20	0.5	10.0	0.40
3	VP1	4	0.2	0.8	10	0.2	2.0	0.40
3	VP2	4	0.5	2.0	10	0.5	5.0	0.40
3	VP3	4	0.7	2.8	10	0.7	7.0	0.40
3	VP4	4	0.8	3.2	10	0.8	8.0	0.40

^a conditions in experiment 1 and 4 were identical; only conditions for experiment 1 are shown here for brevity;

^b the volumes (in microliters) shown are for cohorts 1 and 3. In cohort 2 the volumes were 4.7 instead of 4, 9.4 instead of 10, 14.0 instead of 15, and 20.3 instead of 20 microliters;

^c return rate (expected value);

^d condition BVP1 in experiment 1 was not repeated in experiment 2, but instead the results from experiment 1 were reused in further analyses

Compared to the baselines, mice showed an increase in discrimination performance in the congruent condition and a decrease in performance in the incongruent condition (Fig. 2). Contrary to our expectations, the trade-off between volume and probability did not abolish preference in the incongruent condition (Fig. 1), with a discrimination performance significantly higher than the chance level of 0.5 (lower 95%CI < mean < upper 95%CI, $0.512 < 0.572 < 0.634$). Thus, the volume dimension exerted a stronger influence on choice, at least in absolute terms.

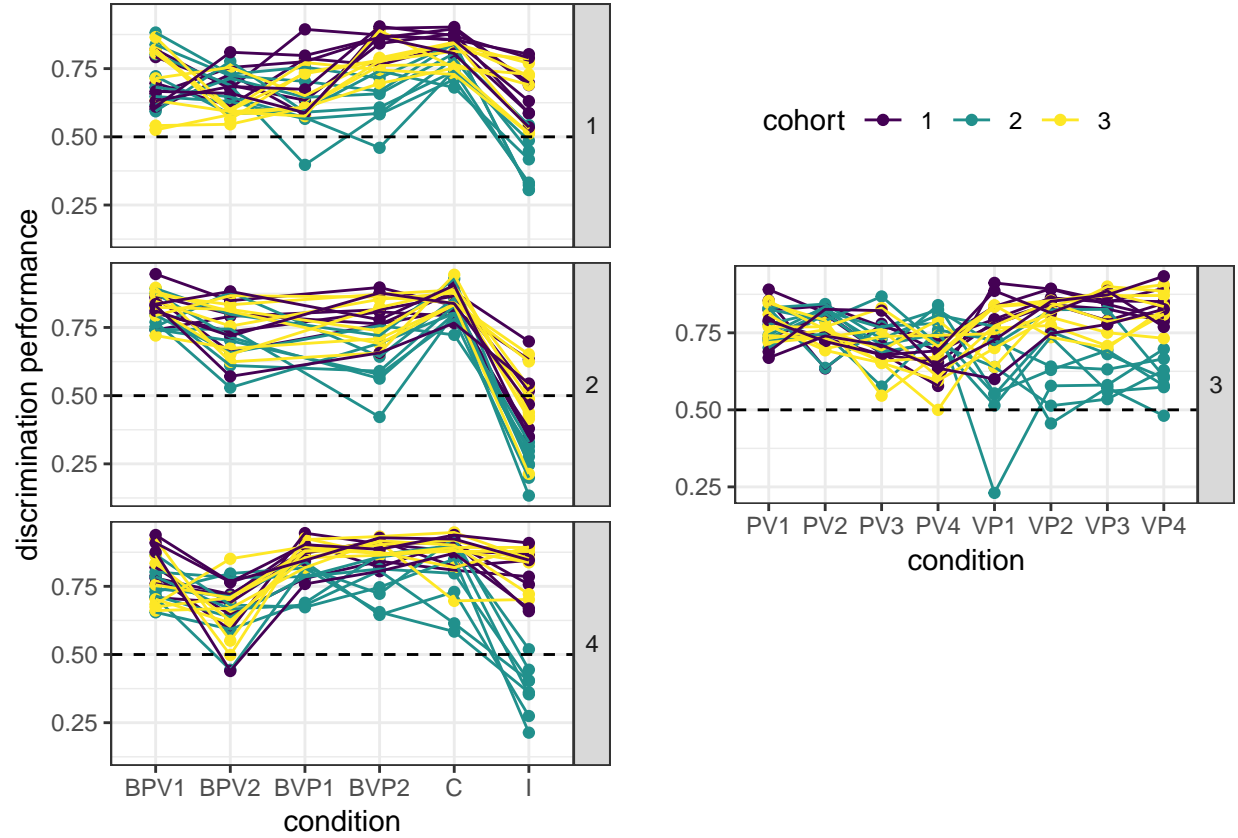


Figure 1: **Overview of discrimination performance for all mice in all experiments.** Experiments 1 through 4 are shown in different panels (1-4). Each symbol is the mean discrimination performance of an individual mouse over two presentations of the same condition (original and reversal). The experimental conditions are described in detail in Table 1. The discrimination performance gives the relative visitation rate of the more profitable option, or, in the incongruent condition, the option with the higher volume. Dashed line gives the chance level of 0.5. Data are shown in different colors for three different cohorts of eight mice each (total $N = 24$). Data from the same individuals are connected with lines. Cohort 2 (green symbols and lines) was tested in a different cage set-up than cohorts 1 and 2 (see Animals, Materials, and Methods for details).

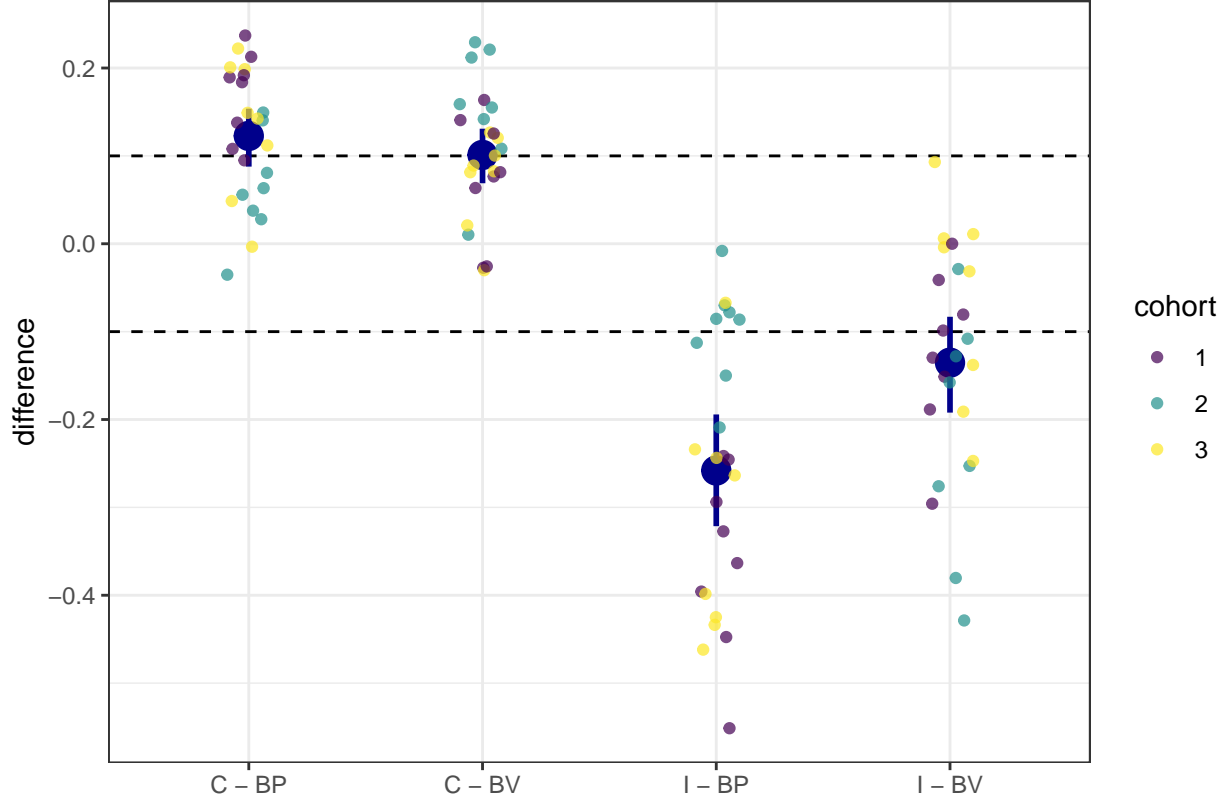


Figure 2: **Difference between discrimination performance in the baseline conditions and in the congruent and incongruent conditions in experiment 1.** Symbols show the individual differences in discrimination performance for the given conditions of each individual mouse ($N = 24$). Mice from different cohorts are shown in different colors. Large blue symbols give the means and the blue vertical lines the 90%-confidence intervals from bootstraps, corrected for multiple comparisons. When the confidence intervals lie completely within the smallest effect size of interest (sesoi) interval bounded by the dashed lines, there is statistical support for equivalence (Lakens 2017). When the confidence intervals do not cross the zero line, there is statistical support for difference. If the confidence intervals cross the zero line, but are not completely bounded by the sesoi, the results are inconclusive. The discrimination performances in the baseline conditions were calculated from the mean values from the two different baseline conditions for each reward dimension (volume and probability), i.e. BP was the mean of BPV1 and BPV2, and BV was the mean of BVP1 and BVP2 (Table 1). The discrimination performance in the incongruent condition was calculated as the relative preference for the higher probability dispenser when contrasted with the probability baseline (I - BP) and for the higher volume dispenser when contrasted with the volume baseline (I - BV).

Experiment 2

In previous experiments (Rivalan, Winter, and Nachev 2017), we had shown that the relative stimulus intensity, i.e. the absolute difference between two options divided by their mean, was a good predictor of discrimination performance for both volume and probability differences. Another finding from these experiments was that, at least initially, mice responded less strongly to differences in volume than to differences in probability, despite equivalence in return rates (Rivalan, Winter, and Nachev 2017). We tried to correct for this in experiment 1 by selecting options with a higher relative intensity for volume (4 μ L vs. 20 μ L, rel.int. = 1.33) than for probability (0.2 vs. 0.5, rel.int. = 0.857). In order to test whether we had over-corrected for decreased sensitivity to volume in experiment 1, we performed a slightly modified version, experiment 2, which was the same, but with probability of 1 instead of 0.5 in every choice option from experiment 1 (Table 1). Thus, with the two choice options having the same relative intensities (rel.int. = 1.33) and being equivalent in return rates, we expected the discrimination performance in the incongruent condition to be at chance level if both dimensions were equally weighed and equally perceived. On the other hand, if mice were less sensitive for volume than for probability differences as in our previous experiments, then the discrimination performance in the incongruent condition should be skewed towards probability (< 0.5).

In contrast to experiment 1, in experiment 2 mice showed an increase in discrimination performance in the congruent condition only when compared to the volume baseline, but not when compared to the probability baseline (Fig. 3). As in experiment 1, the discrimination performance in the incongruent condition was lower than in either of the two baselines (Fig. 3). Although the discrimination performance in the incongruent condition was again different from 0.5 ($0.349 < 0.407 < 0.472$), it was lower than chance, thus skewed towards probability (Fig. 1).

Experiment 3

In the previous experiments we used two different baseline conditions for each dimension (BPV1, BPV2, BVP1, and BVP2, Table 1), in order to exhaust all combinations of reward stimuli and balance the experimental design. But could it be that the level of the background dimension despite being the same across choice options nevertheless affected the discrimination performance on the relevant dimension? Researches have proposed that in multi-dimensional choice the decision process can be considerably simplified if differences that are (nearly) equal are not evaluated but ignored (Tversky 1969; Shafir 1994; Shafir and Yehonatan 2014). Thus we can predict that regardless of the level of the background dimension, the discrimination performance on the relevant dimension should remain constant. Alternatively, animals could use all information from every reward dimensions for the estimation of a single value (utility), in what some authors refer to as “absolute reward evaluation” (Tversky 1969; Shafir 1994; Shafir and Yehonatan 2014). Since the utility curve is generally assumed to progressively increase with the increase in any given good, but with a decreasing slope (Kahneman and Tversky 1979; Kenrick et al. 2009; but see also Kacelnik and Brito e Abreu 1998), we may expect that as the background dimension increases the subjective difference between the options will decrease and the discrimination performance will also decrease as a result (Shafir and Yehonatan 2014). The same prediction can be made if we assume that the strength of preference increases under lean environmental conditions, i.e. at low reward volume or probability (Schuck-Paim, Pompilio, and Kacelnik 2004). In order to test whether the two reward dimensions (volume and probability) interact with each other even when one of them is irrelevant (being the same across choice options), we performed experiment 3.

The conditions in experiment 3 were chosen to be similar to the background conditions in the previous experiments, by having one background and one relevant dimension (Table 1). The relevant dimension always differed between the two options. For the probability dimension, we selected the same values of 0.2 and 0.5 (rel.int. = 0.86), as in the previous experiments. For the volume dimension we selected the values of 4 μ L (4.8 μ L in cohort 2) and 10 μ L (9.6 μ L in cohort 2), because the combination of a higher volume with a probability of 0.8 was expected to result in an insufficient number of visits for analysis (Table 1). Cohort 2 had different reward volumes due to differences in the pumping process (Animals, Materials, and Methods), which also resulted in a lower relative intensity for volume (0.67 instead of 0.86). There were four different levels for each background dimension (volume and probability, Table 1). Each mouse had its own pseudo-random sequence

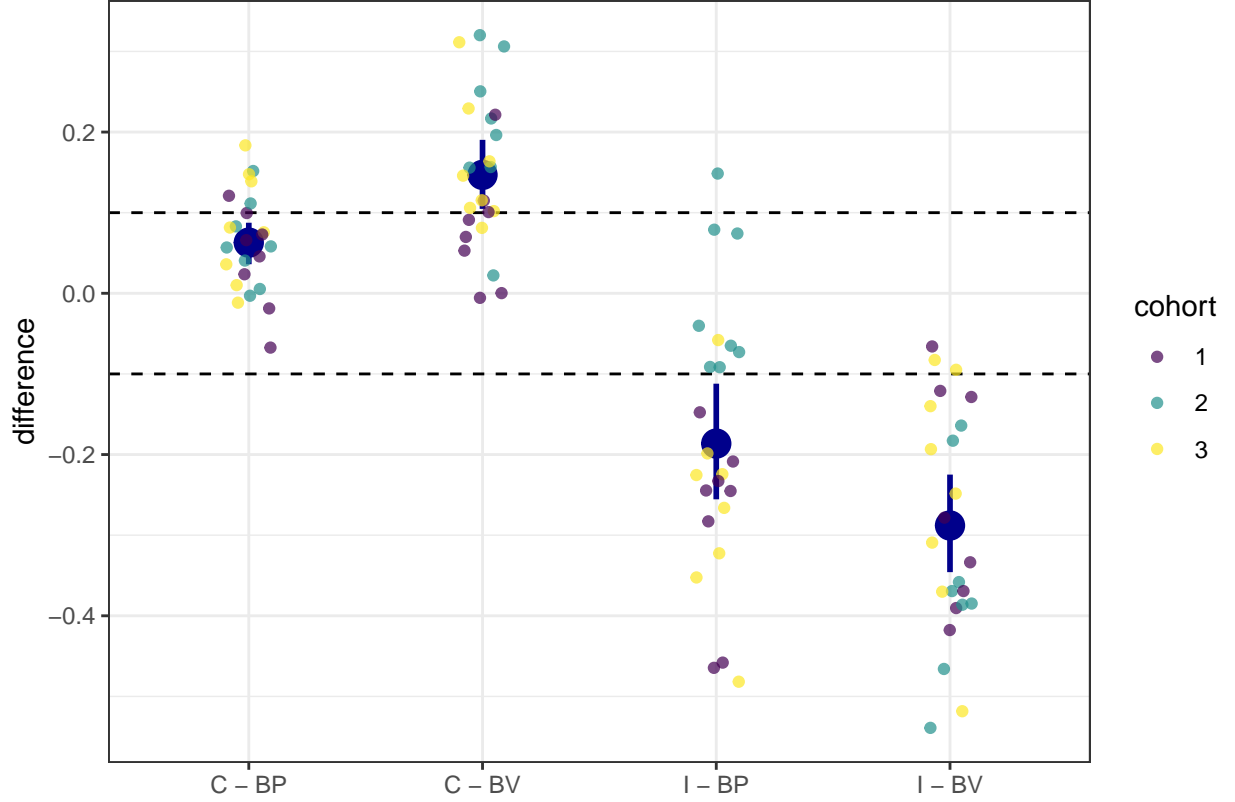


Figure 3: **Difference between discrimination performance in the baseline conditions and in the congruent and incongruent conditions in experiment 2.** Same notation as in Fig. 2. The discrimination performances in the baseline conditions were calculated from the mean values from the two different baseline conditions for each reward dimension (volume and probability), i.e. BP was the mean of BPV1 and BPV2, and BV was the mean of BVP1 and BVP2, where the values for condition BVP1 were taken from experiment 1 (Table 1). The discrimination performance in the incongruent condition was calculated as the relative preference for the higher probability dispenser when contrasted with the probability baseline (I - BP) and for the higher volume dispenser when contrasted with the volume baseline (I - BV).

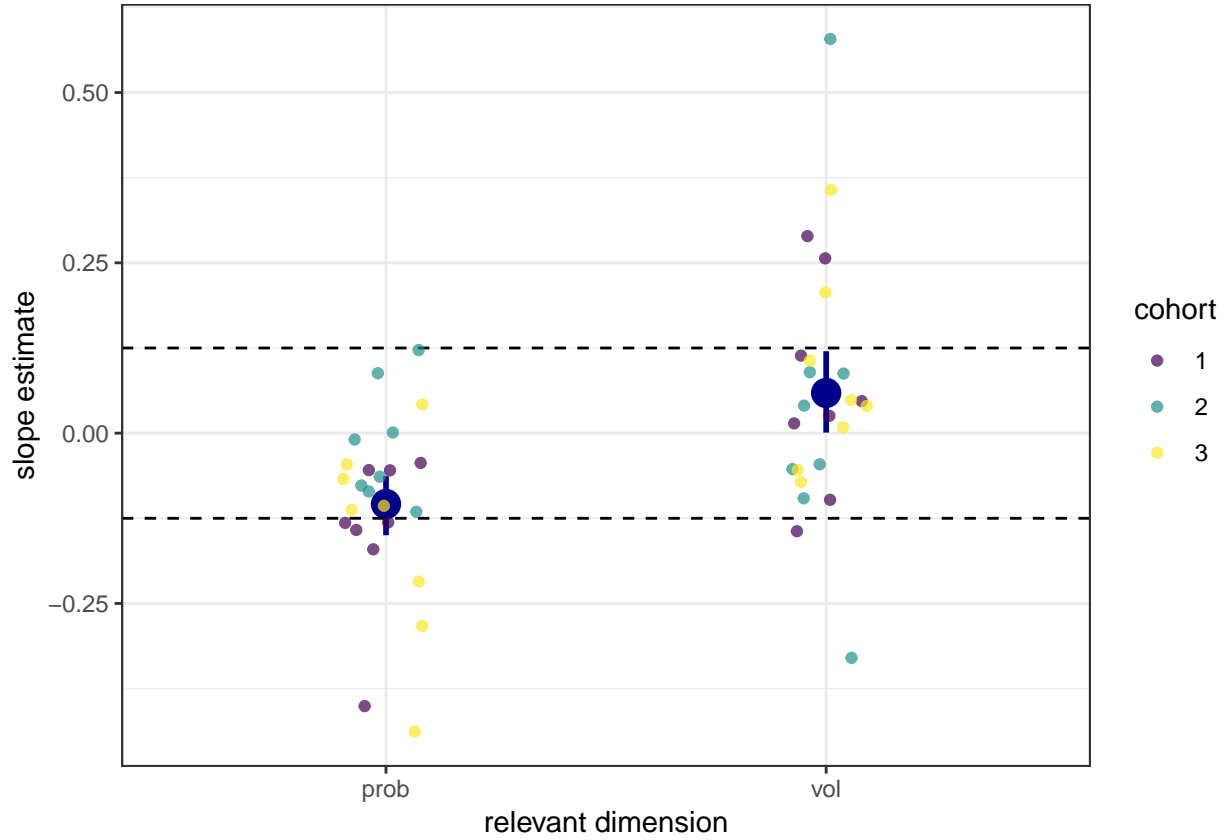


Figure 4: **Slope estimates for the effect of the background dimension on the discrimination performance in the relevant dimension.** The two choice options always differed along the relevant dimension (either probability or volume, given on the abscissa) at a fixed relative intensity. The discrimination performance for each mouse was measured at four different levels of the background dimension, which was set at the same values on both choice options during a single drinking session, but differed from condition to condition (Table 1). Each symbol is the average for an individual mouse over the two presentations of the same condition (original and reversal). The smallest effect size of interest was determined to be the slope that would have resulted in a difference in discrimination performance of 0.1, from the lowest to the highest level of the background dimension. The remaining notation is the same as in Fig. 2.

of the eight possible conditions. As in all other experiments, each condition was followed by a reversal. In order to test whether the background dimension affected discrimination performance, we fitted linear mixed models for each dimension, with discrimination performance as the dependent variable, background level as the independent variable and mouse as a random variable, using lme4 in R (Bates et al. 2015). The background level was the proportion of the actual value to the maximum of the four values tested, e.g. the background levels for volumes 4, 10, 15, 20 were 0.2, 0.5, 0.75, 1, respectively. We defined *a priori* a smallest effect size of interest (sesoi), as 0.125, which is the slope that would result from a difference of 0.1 in discrimination performance between the smallest and the largest background levels (PV1 and PV4, 0.2 and 1, respectively). A slope (whether positive or negative) within the sesoi interval was considered equivalent to zero and demonstrating a lack of an effect of background dimension.

The results of experiment 3 show that the discrimination performance for volume was independent from probability as the background dimension, since the slope was equivalent to zero (Fig. 1, Fig. 4). However, the discrimination performance for probability decreased with increasing volumes, although the effect size was small (Fig. 1, Fig. 4). These results partially support the hypothesis that decision-makers may ignore reward dimensions if options do not vary along them.

Experiment 4

In previous experiments (Rivalan, Winter, and Nachev 2017), mice showed an improved discrimination performance for volume over time. A potential explanation is that, with experience mice become more attuned to the relevant dimension. [More references and explanation on selective attention in multi-dimensional choice here?]

In order to test whether the discrimination performance for one or both dimensions improves over time, we performed experiment 4, which had the same conditions as experiment 1, but with a new pseudo-random order. The same mice participated in all experiments (1-4), with about seven weeks between experiment 1 and experiment 4. We tested the discrimination performance of all mice in each experimental condition for equivalence (Table 1). As in the previous experiments, we also used equivalence tests on the contrasts between the baselines and the congruent and incongruent conditions.

In the comparison between experiment 1 and experiment 4, mice showed an improved discrimination performance in both volume baselines and in the incongruent condition (Fig. 5). There was no change in the BVP2 and C conditions, and the results for BPV1 were inconclusive (confidence interval crosses zero line and not completely bounded by sesoi). Thus, consistent with our prior findings, mice improved their volume discrimination over time. The discrimination performance in the congruent condition was better than in the probability baseline, but the same as in the volume baseline (Fig. 6). The discrimination in the incongruent condition was lower than in any of the two baselines, but the difference to the volume baseline was smaller (Fig. 6). Thus, as in our previous experiments, mice showed an improvement in volume discrimination over time. Furthermore, compared to experiment 1 the influence of the volume dimension on choice was even more pronounced.

Decision models of two-dimensional choice

We based our decision models on the Scalar Utility Theory (SUT, Kacelnik and Brito e Abreu (1998); Rosenström, Wiesner, and Houston (2016)), which models memory traces for reward amounts as normal distributions centered around the remembered value (e.g. volume). The scalar property is implemented by setting the standard deviations of these distributions to be proportional to their means. Choice can then be modeled by taking a single sample from each distribution and selecting the option with the larger sample. As previously explained, the discrimination performance of reward probability can be reasonably predicted by the relative intensity of the two options (Rivalan, Winter, and Nachev 2017). This suggests that the memory traces of reward probabilities or of overall utility also exhibit the scalar property, so that discrimination of small probabilities (e.g. 0.2 vs. 0.5, rel.int. = 0.86) is easier than discrimination of large probabilities (e.g. 0.5

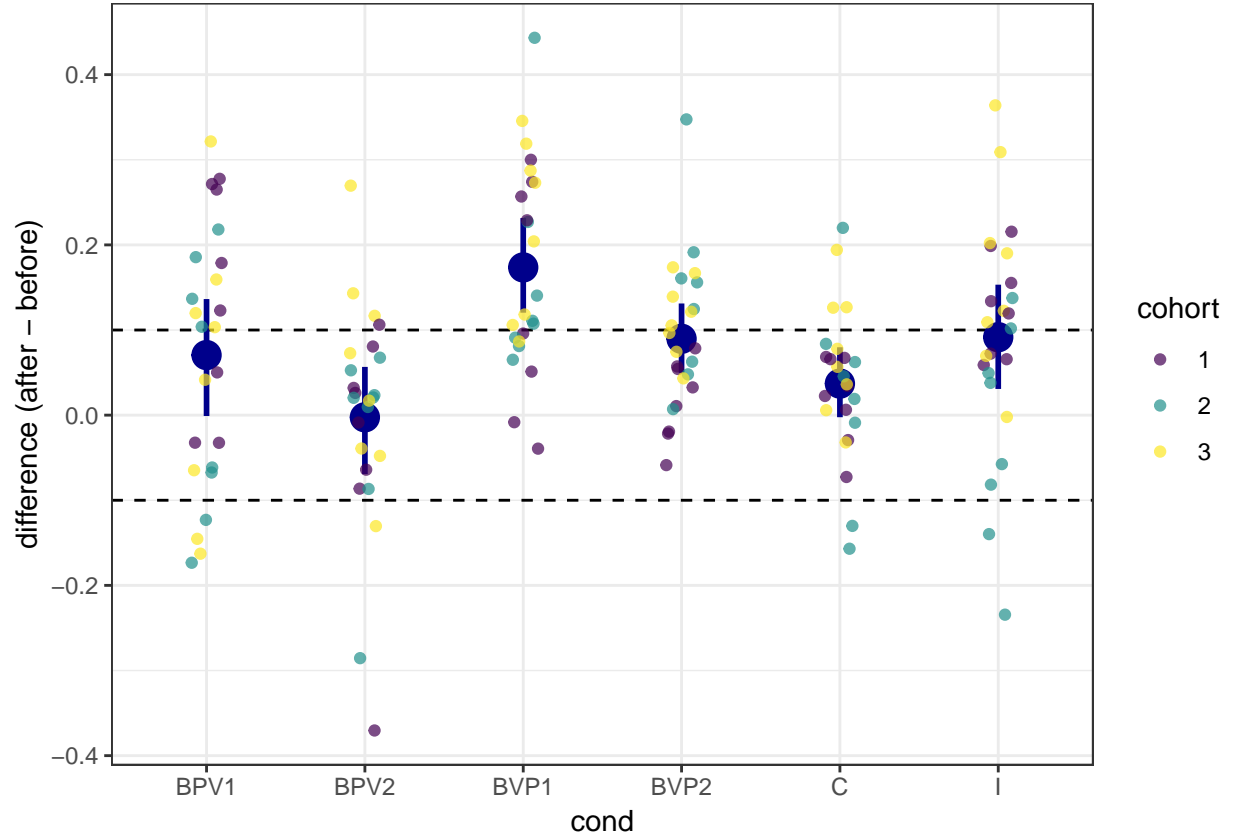


Figure 5: **Difference in discrimination performance between identical conditions in experiment 1 and experiment 4.** Same notation as in in Fig. 2. The sequence of conditions was pseudo-random in each experiment and different for each individual. Positive differences indicate an increase in discrimination performance with time. Mice were seven weeks old at the beginning of experiment 1 and 13-14 weeks old at the beginning of experiment 4. The discrimination performance in the incongruent condition was calculated as the relative preference for the higher volume dispenser.

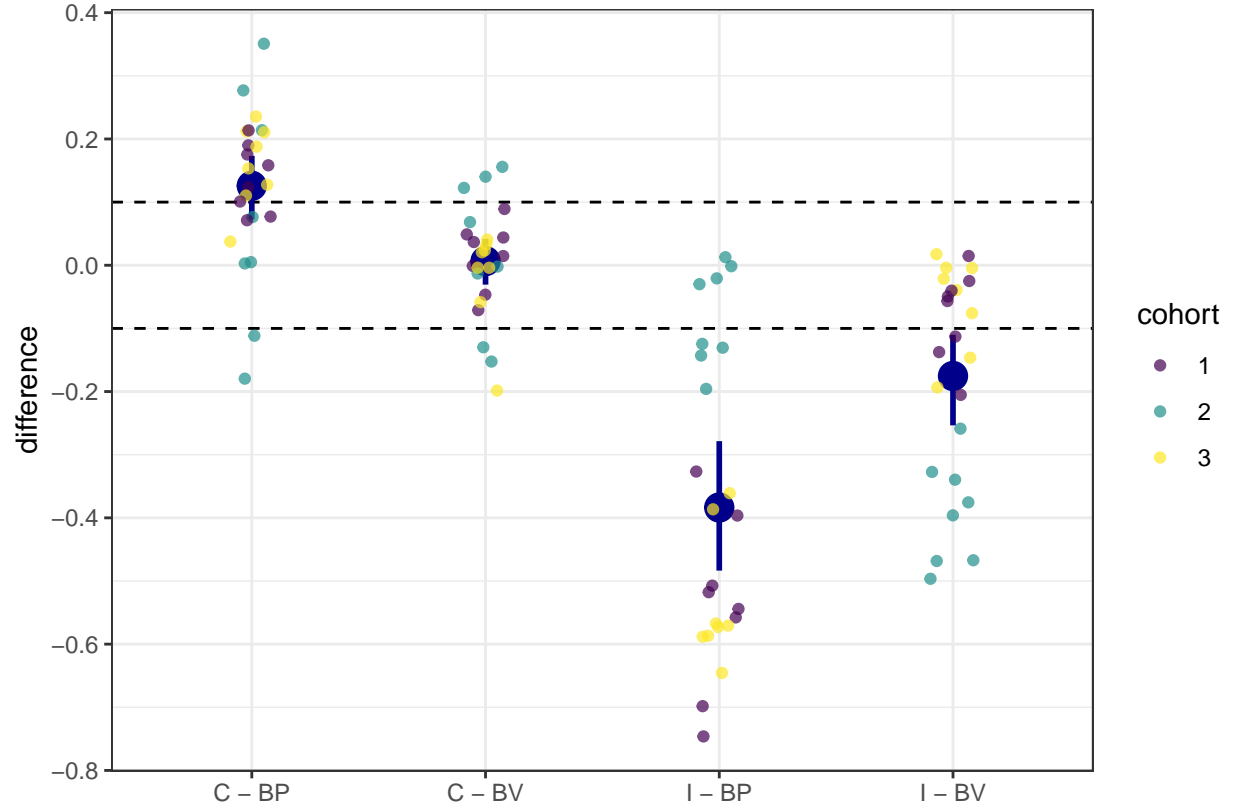


Figure 6: **Difference between discrimination performance in the baseline conditions and in the congruent and incongruent conditions in experiment 4.** Same notation as in Fig. 2. The discrimination performance in the incongruent condition was calculated as the relative preference for the higher probability dispenser when comparing to the probability baseline and for the higher volume dispenser when comparing to the volume baseline. Compare to Fig. 2.

vs. 0.8, rel.int. = 0.46). Consequently, discrimination (of either volumes or probabilities) when options vary along a single dimension can be modeled by SUT.

We next extend this model for multi-dimensional choice situations. We considered various implementations of the base SUT model described above that differed in the integration of information from the volume and probability dimensions. These models were:

1. *Scalar expected value model.* There is a single memory trace for each option and it consists in the simple product of the estimate for the volume and the estimate for the probability (expected value). The scalar property is implemented as $\pi N(v, \gamma \cdot v)$, where v is the volume estimate, π is the probability estimate, and γ is a free parameter, the coefficient of variation. This model thus utilizes information from all dimensions for every decision.
2. *Hurdle model.* There are traces for each dimension for every option, where each trace exhibits the scalar property independently and the value is obtained by simple multiplication of the traces for each dimension: $N(\pi, \gamma \cdot \pi) \times N(v, \gamma \cdot v)$. This model also utilizes information from all dimensions for every decision. Although this model allows each dimension to have its own scalar factor, e.g. $\gamma_\pi \neq \gamma_v$, for the sake of simplicity we assume that they are both equal.

The memory traces in the remaining models are identical to the traces in model 2, but these models usually consider only a single dimension.

3. *Random dimension model.* Each decision is based on a single dimension, selected with probability 0.5.
4. *Winner-takes-all model.* Each decision is based only on the dimension with the higher subjective salience. The salience for a vector of samples (S_i) from the underlying memory traces along one dimension, e.g. volume $v = (S_1, S_2, \dots, S_n)$, is calculated as $n \cdot \frac{\max(v) - \min(v)}{\sum_{i=1}^n v}$, where n is the number of options. In the case of $n = 2$, the salience is equivalent to the previously described relative intensity measure. For dimensions of equal salience the model reverts to model 3.

The last two models are examples of a lexicographic rule, in which the dimensions are checked in a specific order. If the salience of a dimension is higher than a given threshold, then a decision is made based only on this dimension. Otherwise the next-order dimension is checked. If all dimensions have saliences below the threshold, the model reverts to model 3. The value of the threshold was 0.8, the psychometric function threshold for probability (Rivalan, Winter, and Nachev 2017).

5. *Probability first model.* Probability is checked first, then volume.
6. *Volume first model.* Volume is checked first, then probability.

All models described above share the same free parameter, the scalar factor γ . In order to obtain estimates for γ , we first fitted each models to the probability baseline discrimination performances of all mice in experiments 1, and 4 (conditions BPV1 and BPV2). The resulting median estimates (across animals) for γ were then used as the values of the free parameter in out-of-sample tests of the six models. With each model we generated 100 choices by 100 individuals for each experimental condition. We then quantified the model fits to the empirical data by calculating root-mean-square-errors (RMSE), excluding the BPV1 and BPV2 conditions in experiments 1 and 4. Finally, we ranked the models by the RMSE scores. Since we wanted to analyze asymptotic choice, we did not model learning but assumed that the memory traces of the agents were accurate estimates of the physical reward properties.

There was no single model that could best explain the choice of the mice in all experiments, but models 1 (scalar expected value), 2 (hurdle), and 4 (winner-takes-all) were in the top-three performing models for three out of the four experiments (Tables 2, 3, see also Appendix 1 Figures A1, A2, A3, and A4). However, due to the striking differences in performance between cohort 2 and the other cohorts, we also ranked the models separately for the different mouse groups, depending on which cage they performed the experiments in (cohorts 1 and 3 in cage 1 and cohort 2 in cage 2). Indeed, two different patterns emerged for the different cages. For the two cohorts in cage 1, models 1 (scalar expected value) and 4 (winner-takes-all) were the best supported models, followed by 2 (hurdle) and 6 (volume first). Notably, model 6 (volume first) was the best performing model in experiments 3 and 4, but the worst-performing model in experiments 1 and 2. In

Table 2: Decision-making models

model	description
1	scalar expected value
2	hurdle
3	random dimension
4	winner takes it all
5	probability first
6	volume first

Table 3: Best performing models ranked by root-mean-square-errors (RMSE).

rank	experiment			
	1	2	3	4
1	2	2	6	1
2	3	1	1	4
3	4	4	3	2
4	1	3	4	3
5	5	5	2	6
6	6	6	5	5

contrast, model 5 (probability first) was the best supported model for cohort 2 in all experiments, followed by models 3 (random dimension) and 2 (hurdle).

Table 4: Best performing models ranked by root-mean-square-errors (RMSE) for cohorts 1 and 3.

rank	experiment			
	1	2	3	4
1	2	1	6	6
2	1	4	1	1
3	4	2	3	4
4	3	3	4	2
5	5	5	2	3
6	6	6	5	5

Table 5: Best performing models ranked by root-mean-square-errors (RMSE) for cohort 2.

rank	experiment			
	1	2	3	4
1	5	5	5	5
2	3	2	4	3
3	2	3	1	2
4	4	1	3	4
5	1	4	2	1
6	6	6	6	6

198

199
200
201
202
203
204
205
206
207
208
209
210

211
212
213
214
215
216
217
218

219

220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240

241

242

243

244

NA, NA), c(0.12, 0.12) between a volume background at 4 μ L and at 20 μ L). This suggests that the two dimensions interact with each other. Absolute reward evaluation (Shafir 1994; Shafir and Yehonatan 2014) and state-dependent evaluation (Schuck-Paim, Pompilio, and Kacelnik 2004) are both consistent with this decrease in discrimination performance, but not with the lack of effect in the conditions in which the probability was the background dimension. With comparable return rates (Table 1) between the two series of conditions, these hypotheses make the same predictions regardless of which dimension is relevant and which is background. An alternative explanation is that arriving at a good estimate of probability requires a large number of visits and when the rewards are richer (of higher volume), mice satiate earlier and make a smaller total number of visits, resulting in poor estimates of the probabilities and poorer discrimination performance. Consistent with this explanation, mice made on average (\pm SD) 474 ± 199 nose pokes at the relevant dispensers at 4 μ L, but only 306 ± 64 nose pokes at 20 μ L (Fig. A5).

As mentioned earlier, researchers have proposed that with absolute reward evaluation the difference/mean ratio in an experimental series like our experiment 3 should decrease with the increase of the background dimension, leading to a decrease in the proportion preference for the high-profitability alternative (i.e. discrimination performance) (Shafir and Yehonatan 2014). However, if we multiply the estimates for each dimension together, as in our models 1 and 2, we have an implementation that qualifies as absolute reward evaluation (because a single utility is calculated from the different reward dimensions), which however does not satisfy this prediction (Fig. A6). In fact, none of our models exhibited an effect of the background dimension in experiment 3 on the discrimination performance, with all slopes equivalent to zero (Fig. A6). Thus, our results also show that the background dimensions need not have an effect on the discrimination performance, even with absolute reward evaluation.

Difference between cohorts

The most likely explanation that we can give for the unexpected difference in behavior between the cohorts (most obvious in Fig. 6) is a cage effect. As explained in Animals, Methods, and Materials, the precision of the reward volumes was lower in cage 2, which housed cohort 2. It is unlikely that such a small magnitude of the difference ($0.33 \pm 0.03 \mu\text{Lstep}^{-1}$ in cage 1 vs. $1.56 \pm 0.24 \mu\text{Lstep}^{-1}$ in cage 2) could influence volume discrimination to the observed extent. Future experiments can address this issue by specifically manipulating the reliability of the volume dimension using the higher-precision pump. However, we suspect that the difference between cohorts might have been caused by the acoustic noise and vibrations produced by the stepping motors of the pumps. The pump in cage 1 was much louder, whereas the one in cage 2 was barely audible (to a human experimenter). This could have made it harder for mice in the soft cage (cage 2) to discern whether a reward was forthcoming. As a result, mice waited longer before leaving the dispenser during unrewarded nose pokes (Fig. 7). This potentially costly delay might have increased the relative importance of the probability dimension, resulting in the observed discrimination performance in cohort 2. Furthermore, the same line of reasoning can also explain the improved volume discrimination from the first to the fourth experiment: there was a shift towards shorter visit durations in the cohorts in the loud cage (cage 1) from the first to the fourth experiment (Fig. 7), suggesting that mice had learned over time to abort the unrewarded visits. In an unrelated experiment we tested two cohorts of mice in both cages simultaneously and then translocated them to the other cage. The results demonstrated that differences in discrimination performance were primarily influenced by cage and not by cohort (data not shown). Thus, the sound cue associated with reward delivery may be an important confounding factor in probability discrimination in mice, as it provides a signal for the reward outcome (Ojeda, Murphy, and Kacelnik 2018).

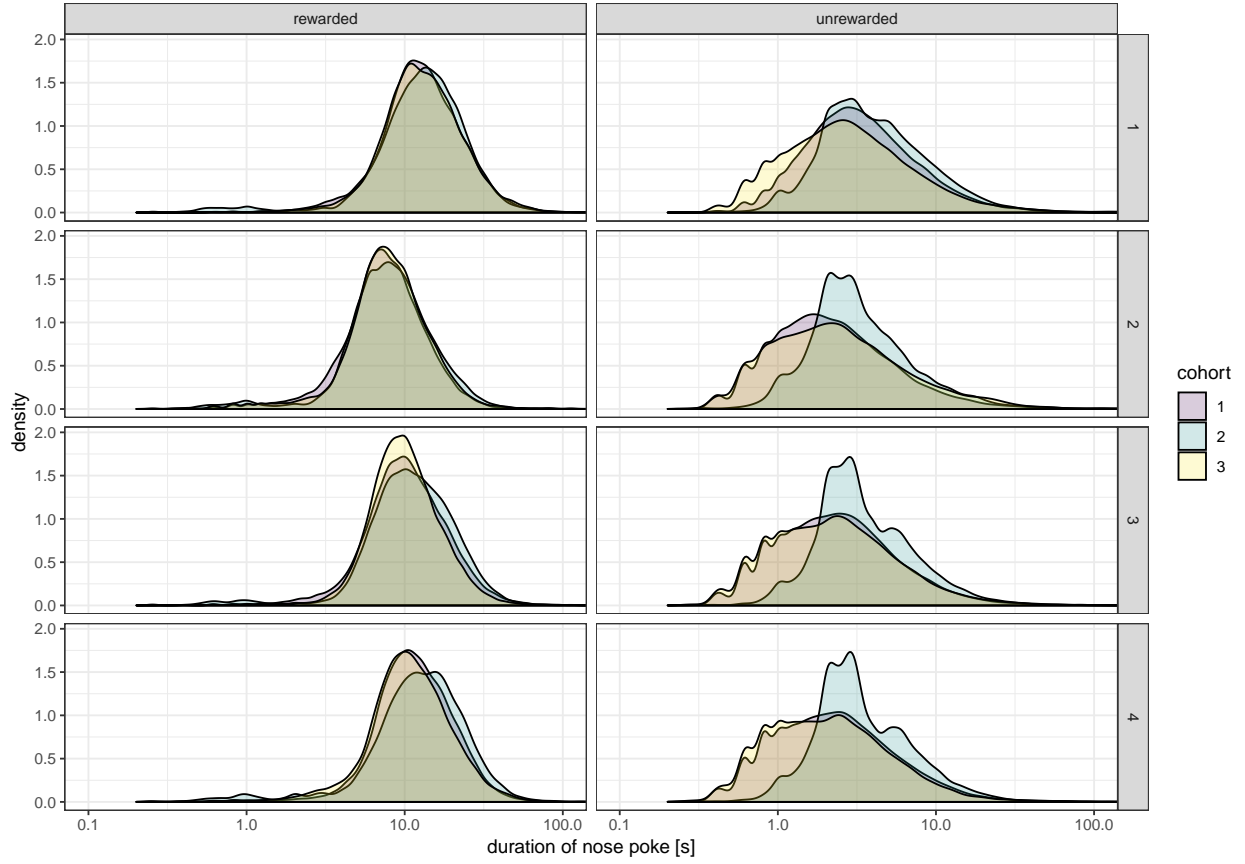


Figure 7: **Visit durations during rewarded and unrewarded nose pokes for the three cohorts in all experiments.** Columns give the status of the nose poke (rewarded or unrewarded) and rows, the experiment number (1-4). Data from the three cohorts are represented by differently color-filled density curves from the observed individual nose poke durations. Note the logarithmic scale on the abscissa.

Animals, Methods, and Materials

Subjects

The experiments were conducted with C57BL/6NCrl mice (Charles River, Sulzfeld, Germany, $N_{\text{total}} = 30$). Mice were five weeks old on arrival. The mice from each cohort were housed together. They were marked with unique Radiofrequency Identification tags (RFID: 12×2.1 mm, 125 kHz, Sokymat, Rastede, Germany) and also earmarked at age six weeks. At age seven weeks mice were transferred to the automated group home cage for the main experiment. Pellet chow (V1535, maintenance food, ssniff, Soest, Germany) was always accessible from a trough in the cage lid. Water was available from the operant modules of the automated group cage, depending on individual reward schedules. Light conditions in the experiments were 12:12 LD and climatic conditions were 23 ± 2 °C and 50–70% humidity.

Ethics statement

The experimental procedures were aimed at maximizing animal welfare. During experiments, mice remained undisturbed in their home cage. Data collection was automated, with animals voluntarily visiting waters dispensers to drink. The water intake and health of the mice was monitored daily. Due to the observational nature of the study, animals were free from damage, pain, and suffering. The animals were not sacrificed at the end of the study, which was performed under the supervision and with the approval of the animal welfare officer (*Tierschutzbeauftragter*) heading the animal welfare committee at the Humboldt University. Experiments followed national regulations in accordance with the European Communities Council Directive 10/63/EU.

Cage and dispenser system

We used automated cages ($612 \times 435 \times 216$ mm, P2000, Tecniplast, Buggugiate, Italy) with woodchip bedding (AB 6, AsBe-wood, Gransee, Germany), and enriched with two grey PVC tubes and paper towels as nesting material. The cage was outfitted with four computer-controlled liquid dispensers. The experimental set-up is described in detail in Rivalan, Winter, and Nachev (2017). Briefly, mice were detected at the dispensers via infrared beam-break sensors and RFID-sensors. Water delivery at each dispenser could be controlled, so that it could be restricted or dispensed at different amounts on an individual basis. Mice were therefore rewarded with droplets of water from the dispenser spout that they could remove by licking. We changed cage bedding and weighed all animals on a weekly basis, always during the light phase and at least an hour before the start of the testing session. Data were recorded and stored automatically on a laptop computer running a custom-written software in C#, based on the.NET framework. Time-stamped nose poke events and amounts of water delivered were recorded for each dispenser, with the corresponding mouse identity.

A second automated group cage (cage 2) was made for the purposes of this study, nearly identical to the one described above (cage 1). The crucial modification was that the stepping-motor syringe pump was replaced with a model that used disposable plastic 25-mL syringes instead of gas-tight Hamilton glass syringes (Series 1025). Thus, the pumping systems in the two cages differed in the smallest reward that could be delivered and in the precision of reward delivery (mean \pm SD: $0.33 \pm 0.03 \mu\text{Lstep}^{-1}$ in cage 1 vs. $1.56 \pm 0.24 \mu\text{Lstep}^{-1}$ in cage 2). The precision of each pump was estimated by manually triggering reward visits at different preset pump steps (17 and 42 in cage 1, 3 and 12 in cage 2) and collecting the expelled liquid in a graduated glass pipette placed horizontally next to the cage. Each dispenser was measured at least 20 times for each pump step value.

Experimental schedule

The different experimental conditions for all animals and cohorts are listed in Table 1.

331 **Pre-training (include discrepancies between cohorts)**

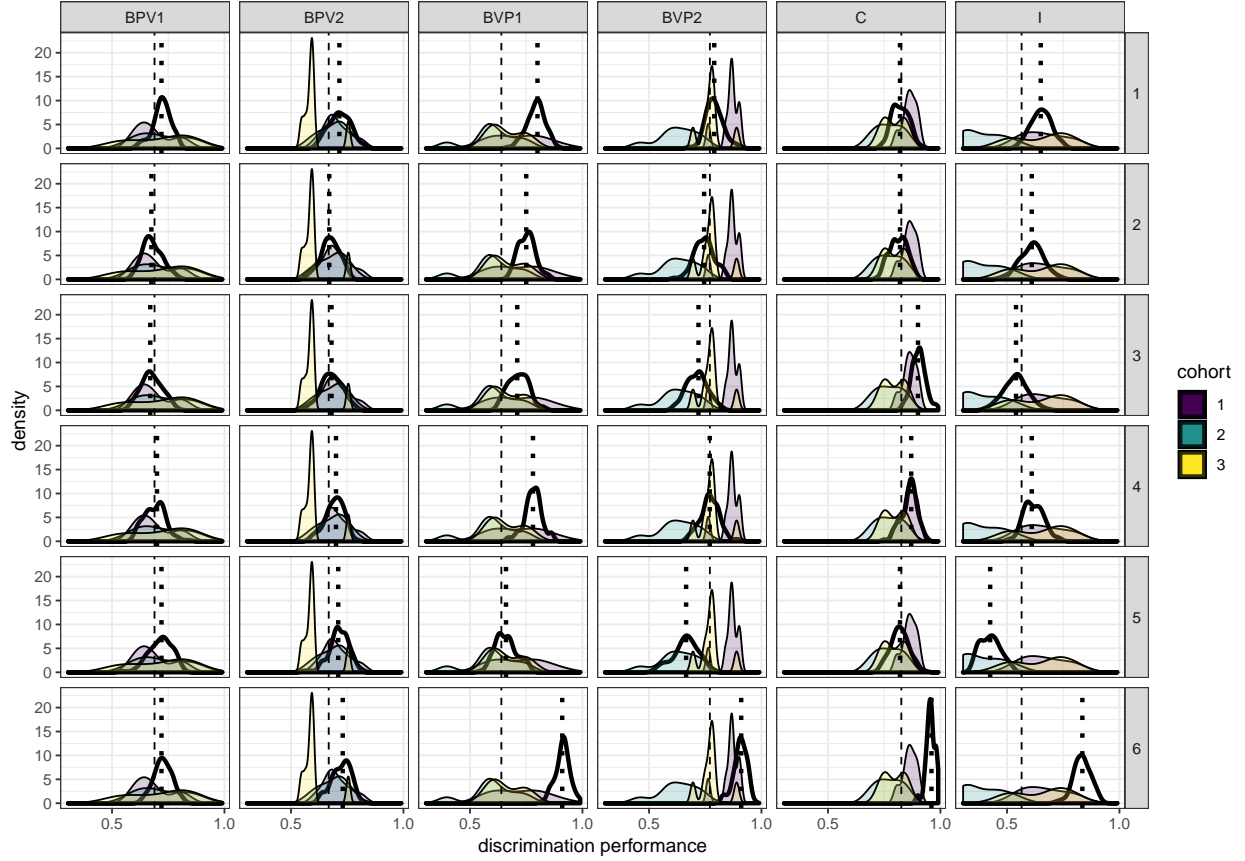


Figure A1: **Comparison of discrimination performances in all six simulation models and in the three mouse cohorts in Experiment 1.** Columns give the condition names (Table 1) and rows, the model number (Table 2). Empirical data from the three cohorts are represented by differently color-filled density curves from the observed discrimination performances. Simulation data are represented by an empty thick-lined density curve. The dashed line gives the median of the empirical data and the dotted line - the median of the simulated data. The discrimination performance gives the relative visitation rate of the more profitable option, or, in the incongruent condition, the option with the higher volume.

Appendix

Acknowledgments

We thank Miléna Brunet for data acquisition, Katja Frei for assistance with the experiments, and Alexej Schatz for software programming.

Authorship and contribution

V.N. conceived the study, analysed the data, and programmed the simulations. Y.W. obtained the funding. V.N., M.R., and Y.W. wrote the paper.

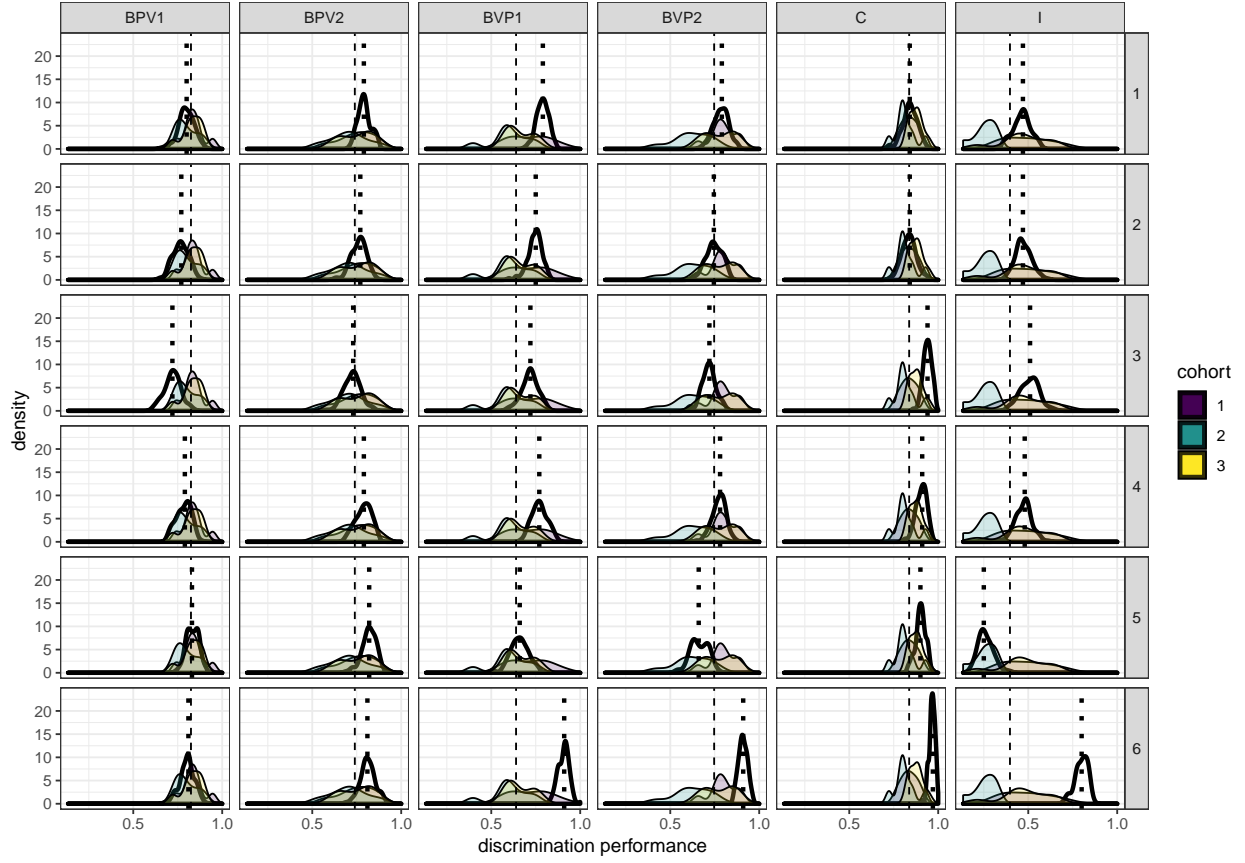


Figure A2: Comparison of discrimination performances in all six simulation models and in the three mouse cohorts in Experiment 2. Same notation as in Fig. A1.

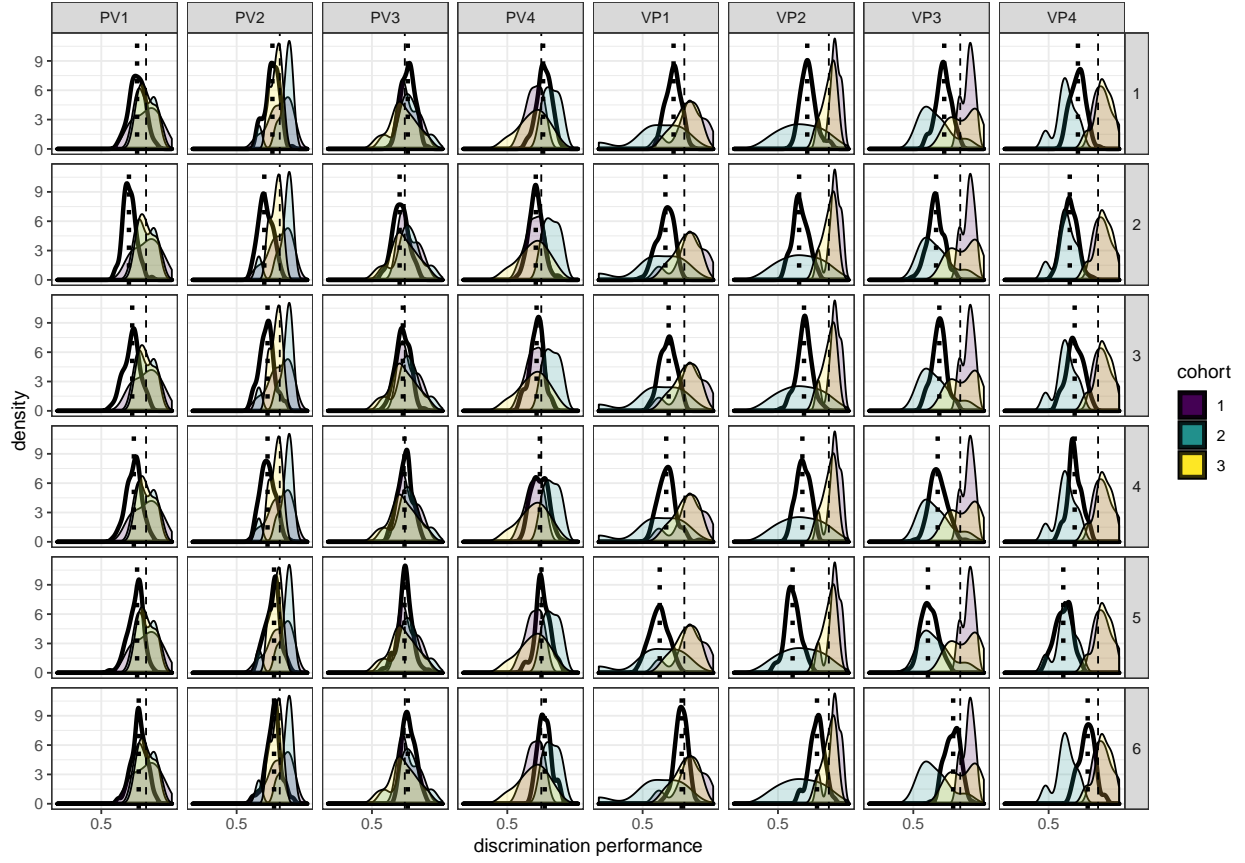


Figure A3: Comparison of discrimination performances in all six simulation models and in the three mouse cohorts in Experiment 3. Same notation as in Fig. A1.

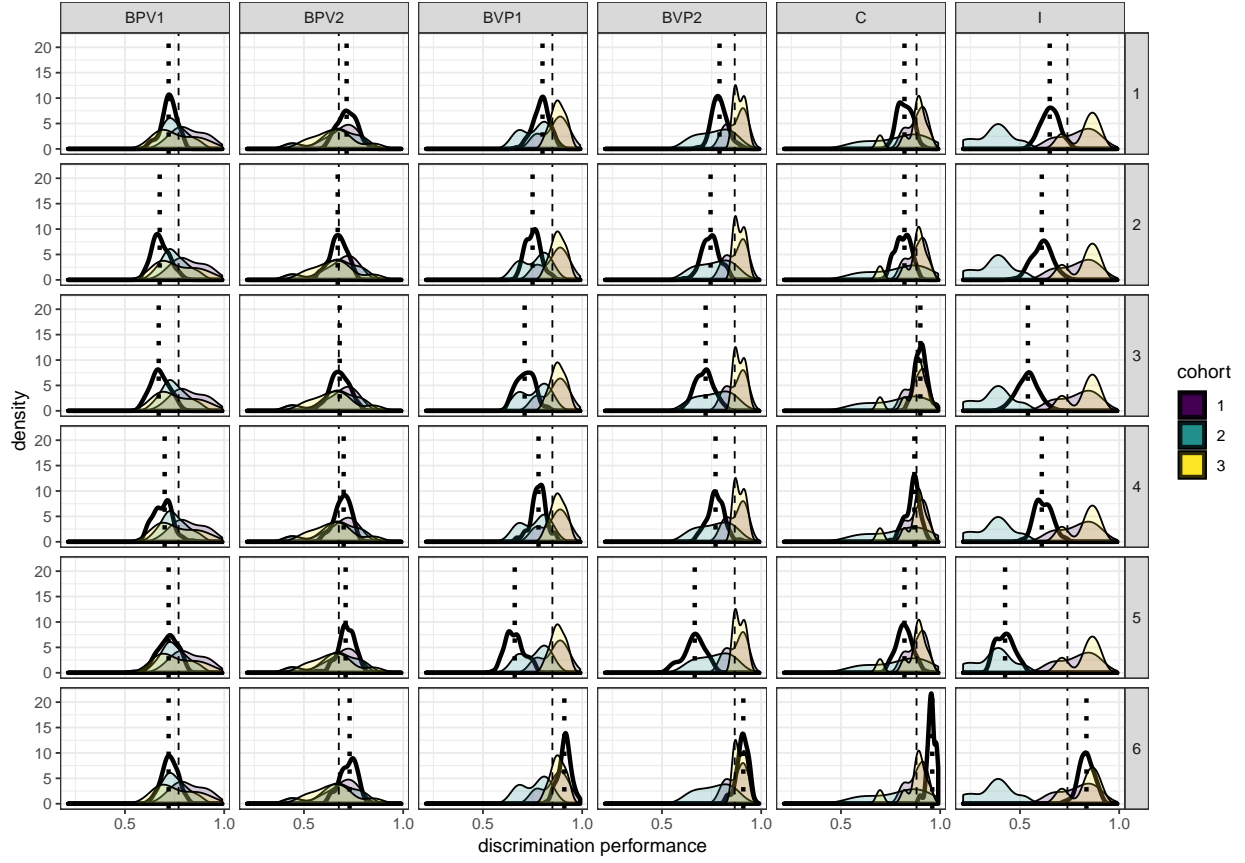


Figure A4: Comparison of discrimination performances in all six simulation models and in the three mouse cohorts in Experiment 4. Same notation as in Fig. A1.

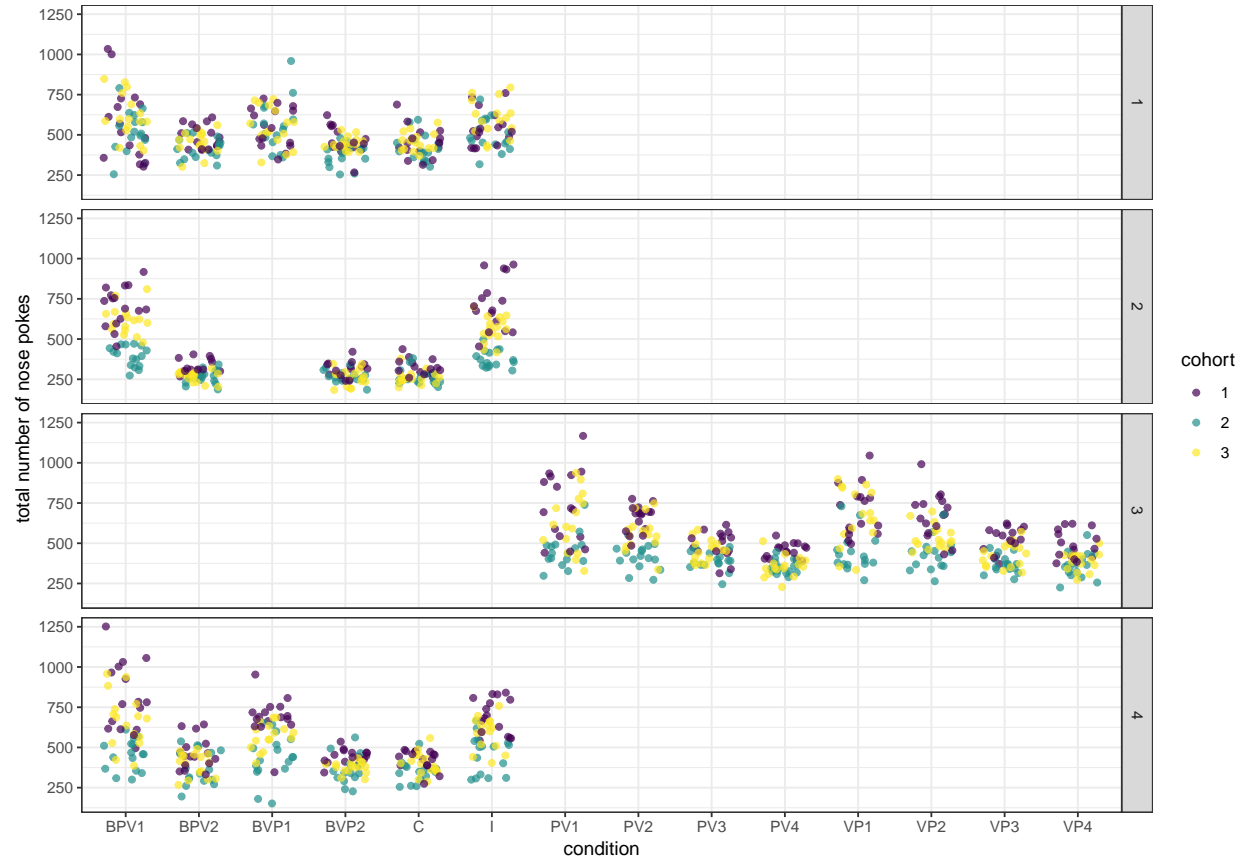


Figure A5: **Total number of nose pokes for each experimental condition in the three cohorts in all experiments.** Rows show different experiments (1-4). Each symbol represents the total number of nose pokes for a single mouse over one of the two experimental days of the given condition.

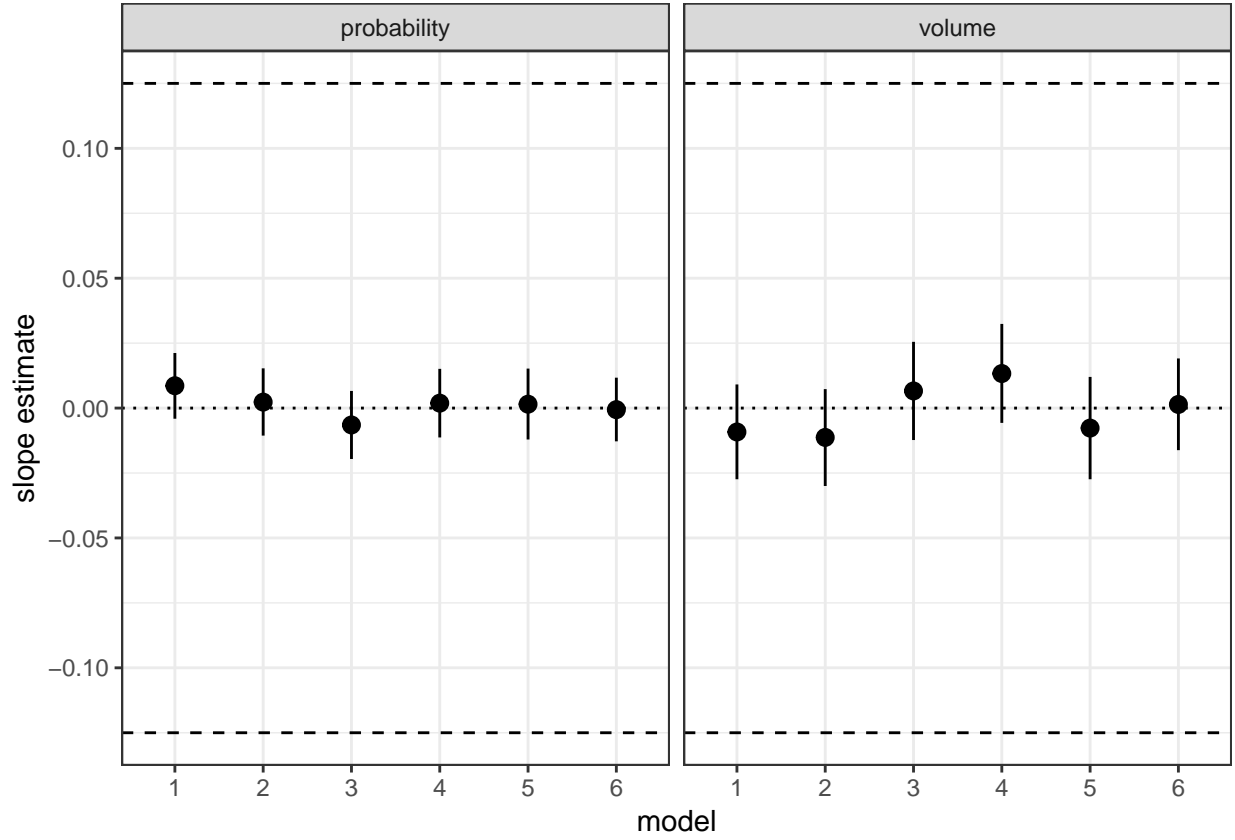


Figure A6: **Slope estimates for the effect of the background dimension on the discrimination performance in the relevant dimension for different decision models.** The two choice options always differed along the relevant dimension (either probability or volume) at a fixed relative intensity. The discrimination performance for 100 virtual mice making 100 decisions each was measured at four different levels of the background dimension. Symbols and whiskers give means and 95% confidence intervals estimated from bootstraps. The smallest effect size of interest (dashed lines) was determined to be the slope that would have resulted in a difference in discrimination performance of 0.1, from the lowest to the highest level of the background dimension. Compare to Fig. 4.

Competing interests

The authors declare that they have no competing interests.

References

- Bates, Douglas, Martin Mächler, Ben Bolker, and Steve Walker. 2015. “Fitting Linear Mixed-Effects Models Using Lme4.” *Journal of Statistical Software* 67 (1):1–48. <https://doi.org/10.18637/jss.v067.i01>.
- Kacelnik, Alejandro. 1984. “Central Place Foraging in Starlings (*Sturnus Vulgaris*). I. Patch Residence Time.” *J Anim Ecol* 53 (1):283–99. <https://doi.org/10.2307/4357>.
- Kacelnik, Alex, and Fausto Brito e Abreu. 1998. “Risky Choice and Weber’s Law.” *J. Theor. Biol.* 194 (2):289–98. <https://doi.org/10.1006/jtbi.1998.0763>.
- Kahneman, Daniel, and Amos Tversky. 1979. “Prospect Theory: An Analysis of Decision Under Risk.” *Econometrica* 47 (2):263–91. <https://doi.org/10.2307/1914185>.
- Kenrick, Douglas T., Vladas Griskevicius, Jill M. Sundie, Norman P. Li, Yexin Jessica Li, and Steven L. Neuberg. 2009. “Deep Rationality: The Evolutionary Economics of Decision Making.” *Soc. Cogn.* 27 (5):764–85. <https://doi.org/10.1521/soco.2009.27.5.764>.
- Lakens, Daniël. 2017. “Equivalence Tests: A Practical Primer for T Tests, Correlations, and Meta-Analyses.” *Social Psychological and Personality Science*, May. <https://doi.org/10.1177/1948550617697177>.
- Levy, Dino J, and Paul W Glimcher. 2012. “The Root of All Value: A Neural Common Currency for Choice.” *Current Opinion in Neurobiology*, Decision making, 22 (6):1027–38. <https://doi.org/10.1016/j.conb.2012.06.001>.
- Matell, Matthew S., Jung S. Kim, and Loryn Hartshorne. 2014. “Timing in a Variable Interval Procedure: Evidence for a Memory Singularity.” *Behav Processes* 101 (January):49–57. <https://doi.org/10.1016/j.beproc.2013.08.010>.
- Nachev, Vladislav, and York Winter. 2019. “Behavioral Repeatability and Choice Performance in Wild Free-Flying Nectarivorous Bats (*Glossophaga Commissarisi*).” *Behav Ecol Sociobiol* 73 (2):24. <https://doi.org/10.1007/s00265-019-2637-4>.
- Ojeda, Andrés, Robin A. Murphy, and Alex Kacelnik. 2018. “Paradoxical Choice in Rats: Subjective Valuation and Mechanism of Choice.” *Behavioural Processes*, Quantitative and comparative analyses of behavior, 152 (July):73–80. <https://doi.org/10.1016/j.beproc.2018.03.024>.
- Rivalan, Marion, York Winter, and Vladislav Nachev. 2017. “Principles of Economic Rationality in Mice.” *Scientific Reports* 7 (1):17441. <https://doi.org/10.1038/s41598-017-17747-7>.
- Rosenström, Tom, Karoline Wiesner, and Alasdair I. Houston. 2016. “Scalar Utility Theory and Proportional Processing: What Does It Actually Imply?” *Journal of Theoretical Biology* 404 (September):222–35. <https://doi.org/10.1016/j.jtbi.2016.06.003>.
- Schuck-Paim, Cynthia, Lorena Pompilio, and Alex Kacelnik. 2004. “State-Dependent Decisions Cause Apparent Violations of Rationality in Animal Choice.” *PLOS Biology* 2 (12):e402. <https://doi.org/10.1371/journal.pbio.0020402>.
- Shafir, Sharoni. 1994. “Intransitivity of Preferences in Honey Bees: Support for ‘Comparative’ Evaluation of Foraging Options.” *Anim. Behav.* 48 (1):55–67. <https://doi.org/10.1006/anbe.1994.1211>.
- Shafir, Sharoni, and Lia Yehonatan. 2014. “Comparative Evaluations of Reward Dimensions in Honey Bees: Evidence from Two-Alternative Forced Choice Proboscis-Extension Conditioning.” *Anim Cogn* 17 (3):633–44. <https://doi.org/10.1007/s10071-013-0694-z>.

- 380 Sih, Andrew, and Marco Del Giudice. 2012. "Linking Behavioural Syndromes and Cognition: A Behavioural
381 Ecology Perspective." *Philos Trans R Soc Lond B Biol Sci* 367 (1603):2762–72. [https://doi.org/10.1098/rstb.](https://doi.org/10.1098/rstb.2012.0216)
382 2012.0216.
- 383 Tversky, Amos. 1969. "Intransitivity of Preferences." *Psychological Review* 76 (1):31–48. [https://doi.org/10.](https://doi.org/10.1037/h0026750)
384 1037/h0026750.