Title:

A passive computational marker for individual differences in non-reinforced learning

Running Title

Computational marker of learning

List of Authors

Tom Salomon[1], Alon Itzkovitch[1], Nathaniel D. Daw[2], Tom Schonberg*[1,3]

[1] Department of Neurobiology, Tel Aviv University.

[2] Neuroscience Institute, Department of Psychology, Princeton University.

[3] Sagol School of Neuroscience, Tel Aviv University.

* Correspondence concerning this article should be addressed to Tom Schonberg, Department of Neurobiology, Faculty of Life Sciences and Sagol School of Neuroscience, Tel Aviv University, Ramat Aviv 6997801, Tel Aviv, Israel. Email: Schonberg@tauex.tau.ac.il.

Conflict of interest

The authors declare no competing financial interests.

Acknowledgements

This work was supported by the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation programme (grant agreement n° 715016), and the Israeli Science Foundation (1798/15 and 1996/20) granted to Tom Schonberg. Tom Salomon was supported by the Nehemia Levtzion fellowship and the Fields-Rayant Minducate Learning Innovation Research Center.

Keywords

Choice Behavior, Decision Making, Learning, Bayesian modeling, individual differences

Abstract

Although research about preference formation and modification has classically focused on the role of external reinforcements, there is also increasing evidence for a key role of non-externally reinforced cognitive mechanisms such as attention and memory in preference modification. In a novel paradigm for behavioral change called the Cue-Approach training (CAT) task, preferences are modified via the mere association of images of stimuli with a neutral cue and a rapid motor response, without external reinforcements. The procedure's efficacy has been replicated across dozens of studies, and the net behavioral change was linked with increased activity in a frontal value-based decision-making brain region during the post-training probe choice phase. However, the cognitive mechanisms during the training phase itself have not been elucidated. Based on the structure of the task alongside recent findings of the involvement of striatal and supplementary motor regions during training, we hypothesized that a motor-related learning process could be a prospective candidate. To test this hypothesis, we developed a computational model of the motor response pattern during training in a large corpus of data collected from 864 participants across 29 different CAT experiments. Using Bayesian modeling of the meta-analysis data, we developed a computational marker for individualized learning in the training task, which was found to be associated with the preference modification effect in the subsequent probe task, both at the participant-level as well as in the more granular individual-item level. Following the conclusions of the meta-analysis, in two additional experiments (a pilot study and a larger preregistered replication study) we aimed to affect learning efficacy by manipulating the training procedure difficulty. As hypothesized and preregistered, training difficulty was captured by the new computational marker identified on the previously collected samples. Manipulation of the training difficulty also resulted in a differential preference modification effect, suggesting a causal relationship between the motor learning captured by the computational model and the post-training behavioral change effect. Our work highlights a novel non-reinforced preference modification pathway, suggesting that attention and motor learning are linked to preference formation, and provides a computational framework to identify individualized training markers which could predict future behavioral change effects.

Introduction

Value construction and modification are commonly understood as a result of learning processes that rely on external reinforcements. In conditioning with external reinforcements an association is formed by associating either a positive or a negative outcome, that lead respectively to enhanced or reduced preference for conditioned stimuli [1–4]. While reinforcement-based learning dominated the research field of value-modifications, a recent review [5] highlights the various means to influence preferences and choices without external reinforcements covered in the literature, dating back to the *mere exposure effect*. In these studies preferences for stimuli can be enhance merely by repeated exposure to stimuli [6]. Similarly, another non-reinforced preference change paradigm that does not rely on external reinforcements can be found in work showing preference modification following previous choices [7]. These studies provide a strong theoretical framing for efficacy of interventions such as advertisements on preferences and how life choices affect future decisions [8,9], above and beyond reinforcement-based manipulations [10,11].

In recent years, a novel procedure named the Cue-Approach Training (CAT) [12] task has been proposed as another means to change preferences without external reinforcements.. The first studies with CAT procedure, showed that preferences for snack food stimuli could be modified via association with a neutral cue and rapid motor response [12–14]. The CAT procedure with snacks included a Go/NoGo training task, in which set of snack food stimuli were associated with a Go cue to which participants were required to respond with a rapid button press response (Go stimuli), while a larger set of snack food stimuli were passively presented for the same exposure period, without a cue or response (NoGo stimuli). The training included several runs in which participants could learn the association of Go stimuli with the cue and response. In a post-training probe phase, when presented with a choice between two snacks for actual consumption, participants demonstrated enhanced preferences for the Go stimuli over NoGo stimuli of similar initial value [12–14].

Subsequent studies with the CAT procedure exemplified its efficacy in modifying preferences for a wide range of stimuli beyond snack food, including healthy food items, faces, positive affective stimuli and fractal art stimuli [15–18]. Overall, across dozens of studies with different types of stimuli, participants consistently demonstrated enhanced preference for Go stimuli over NoGo stimuli of similar initial value [12–20]. Studies that examined the long term maintenance of the CAT effect in additional follow-up sessions, found that the preference modification effect persisted for months without any additional training sessions [12,15,18,21]. The fact

that a preference modification effect was observed following a training procedure with no external reinforcement or feedback, inspired the idea that CAT relies on a non-reinforced valuation pathway, putatively involving attention and motor neural circuits [5].

Several mechanisms have been suggested to underlie non-externally reinforced paradigms. First, the preferences-as-memory theoretical framework suggests that value could actually be framed as retrieval of relevant knowledge about alternatives from memory [22]. This hypothesis was supported by a study showing that Go stimuli that were better remembered were also more preferred [19]. Attention also play plays a key role in value-based decision making both on its own and in synergy with memory [23,24].Support for the involvement of attention was found in eye-tracking studies with CAT, which found that during the probe phase, participants view Go stimuli more [12,13,18], suggesting that increased gaze time during the probe phase indicated attentional evidence-gathering process which increased the likelihood of choosing the Go stimuli over NoGo stimuli [25–27].

While CAT was found to be a reliable means to modify choice behavior, previous studies have mainly focused on the probe phase, leaving ambiguity as to the cognitive mechanisms unfolding during the unique training procedure: specifically, what mechanisms in the training phase itself induce the observed behavioral change. During training all stimuli are presented for the same amount of time, thus mere exposure or viewing time alone cannot account for the preference modification effect. Studies that examined the crucial features of the training task discovered that responding to the Go stimuli also was not sufficient to induce preference changes, as preference modification did not occur in training version in which the Go cue was presented immediately with the Go stimulus, or training where the Go cue was given prior to a block of Go stimuli [13]. These findings suggested that simple attention or motor response are not sufficient, as the rapid motor response during CAT and the motor preparation learning are crucial factors for non-reinforced preference modification with CAT. This conclusion corresponds with a recent imaging study with CAT [18], which found that individual differences in the preference modification effect correlated with increased neural activity during training within the supplementary-motor cortex and the striatum, regions well associated with motor-planning [28,29] and reinforcement-based learning [30–33], respectively. The striatum has also been suggested to be key neural hub for learning thanks to its unique dual-role in both reward processing and motor regulation networks [34,35].

Based on these combined results, we sought to utilize the large corpus of data collected with the CAT task to identify the underlying mechanism of non-reinforced preference modification, with emphasis of the training phase. An important aspect of the CAT is that training is performed at the individual item basis, and preference change is tested in a subsequent binary choice phase. Therefore, we surmised that identifying individual differences in motor learning during the training phase could serve as a key factor to explain differential non-reinforced behavioral change effect following CAT. We aimed to test this hypothesis by developing an individualized learning marker based on training motor-response data and test its association with the preference modification effect observed in the subsequent probe phase. To achieve this aim we first developed a novel computational marker using Bayesian modeling of reaction-time (RT) patterns during CAT. Then, based on this model we designed a new non-reinforced training procedure to test our hypotheses directly.

The current work thus consists of two main studies. The first part includes a meta-analysis study, where using data of 864 participants from 29 previous CAT experiments, we devised a novel marker for learning. By combining multiple samples with the CAT into one meta-analysis, we were able to create a unique dataset of a standardized non-reinforced preference modification task. To identify an individualized marker for learning, we first examined RT patterns during the training phase in an exploratory analysis and built a computational model of these patterns using a Bayesian computational framework which included modeling an individualized learning parameter. We examined the association of the individualized learning marker with individual differences in preference modification, as measured in the subsequent binary-choice probe phase. We further examined the hypothesis that the mechanism of non-reinforced preference modification is operating both at the participants-level (i.e., some participants learn better than others), as well as on a more granular item-level (i.e., within participants, some stimuli are better learned than other). We expanded our computational model to capture variability in within-participant item-level learning and examined its association with subsequent choice pattern.

In the second part, we implemented the findings from the Bayesian model that was based on the meta-analysis to develop a novel CAT design and directly manipulate the calculated parameter found in the previous part. In the new design, we used two different cue-contingency conditions to manipulate learning difficulty. We hypothesized that the cue contingency manipulation will affect the motor response and will be captured by the individualized learning

marker devised in the first study, as well as predict subsequent preference modification in the probe phase. The second phase included one preliminary experiment (n = 20), which was followed by a larger pre-registered replication study (n = 59). All the hypotheses, experimental design and analysis plans were carefully preregistered before data collection for the replication study began (https://osf.io/nwr4v). By manipulating the task design to induce differential behavioral modification effect, we aimed to establish a causal relationship between the proposed motor cognitive mechanism and the preference modification effect.

Developing an individualized computational marker for CAT would shed light on the mechanisms underlying successful non-reinforced preference modification and would allow to identify learning patterns both between-participants and within-participant, at the individual item level. These patterns could potentially enable monitoring learning in real-time and the development of novel closed-loop interventions.

Study 1: Meta-analysis of CAT studies

The first part of the current work was a meta-analysis of 29 CAT experiments with a total of N = 864 participants, conducted by our research group and colleagues (see methods for sample size, publication origin and demographics of the individual experiments). All experiments included the three main phases of the CAT procedure: initial preference evaluation, cue-approach training, and a preference modification probe task (Figure 1; see detailed description in the Methods section). In the first phase of the task, initial baseline preferences for a set of stimuli were evaluated in an auction procedure [36], when snacks were used, or a forced choice task [15] for all other stimuli types . Following initial preference evaluation, in the cue-approach training phase, approximately 30% of the stimuli were consistently associated with a neutral Go cue (Go stimuli), to which participants needed to respond with a rapid button press, while the rest of the stimuli were presented passively without cue or response (NoGo stimuli). Participants were instructed to wait for the Go cue and respond as fast as they could, once they perceived the cue. In the final probe phase, preference modification following CAT was evaluated. Participants were asked to indicate their preferred stimulus out of pairs of stimuli with similar initial value, in which one of the two stimuli was a Go stimulus and the other was a NoGo stimulus.

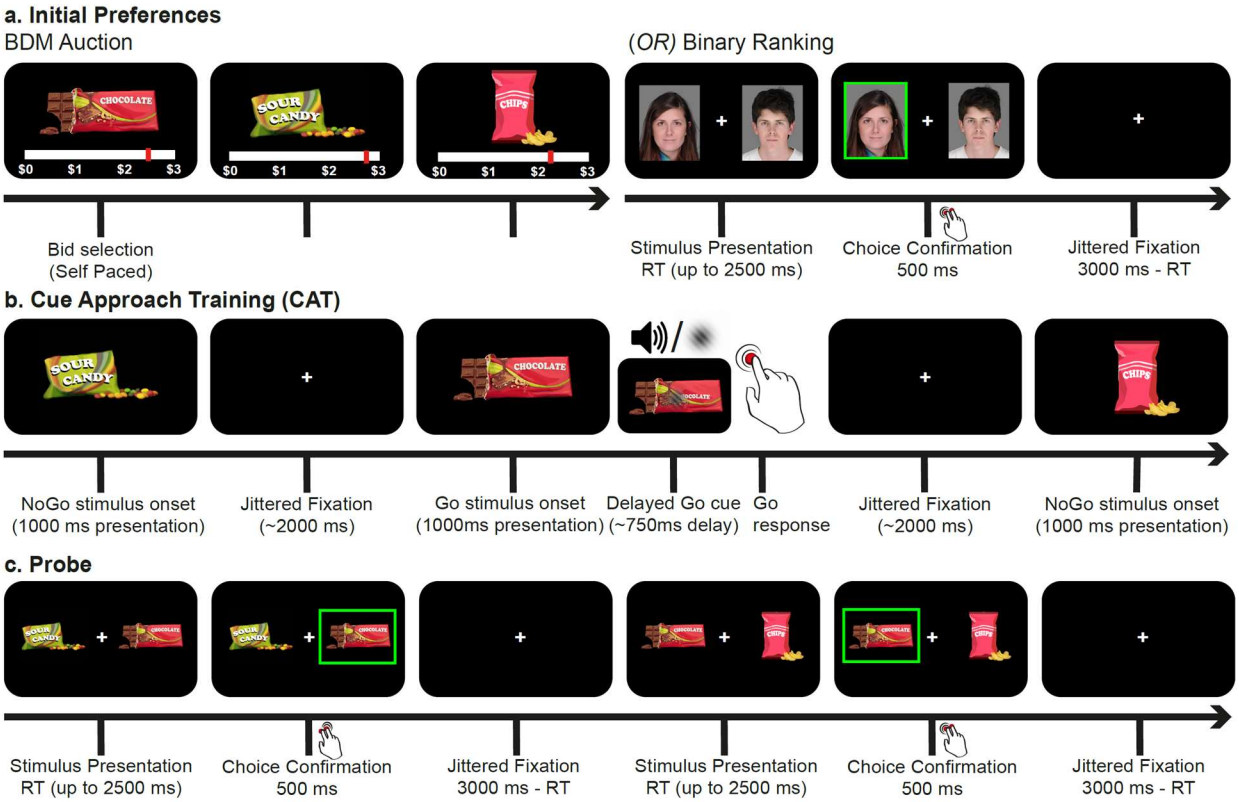


Figure 1. General outline of the three main procedural components of the Cue-approach training (CAT) paradigm. (a) Initial preference evaluation task. Baseline preferences for all stimuli were evaluated either with using a Becker-DeGroot-Marschak (BDM) auction (for consumable stimuli) or using a forced choice binary ranking task (for non-consumable stimuli). (b) In the cue approach training (CAT) task, approximately 30% of stimuli were presented in association with a delayed cue (auditory or visual) to which participants responded with a rapid button press (Go stimuli). All other stimuli were presented without cue and response (NoGo stimuli). (c) In the probe phase, preference modification was evaluated using a binary forced choice between pairs of stimuli of similar initial value, where one was a Go stimulus and the other a NoGo stimulus. Face images are included with permission from the copyright holder [37].

The meta-analysis study focused on identifying response patterns in the training task which could indicate learning efficacy and predict the behavioral change measured in the subsequent probe task. We hypothesized that faster RT could indicate improvement in learning.

Results

RT analysis of CAT. The training task in the different experiments consisted of 8-20 training runs (the number of training run varied between experiments and identical within each experiment). During each run, all Go and NoGo stimuli were presented once individually on-screen. During NoGo trials, stimuli were presented without a Go cue and required no response. Whenever Go stimulus appeared (approximately 30% of trials), a Go cue was followed, to which

200 participants were asked to respond with a rapid button press, before stimuli offset. Participants
201 were instructed to respond *only* when they hear or see the cue. Each trial started with stimulus
202 onset, which was presented on screen for 1000ms. During Go trials, a Go cue appeared
203 approximately 750ms following stimulus onset (thus leaving participants approximately 250ms to
204 respond). The cue onset changed according to participants' performance, so that each successful
205 Go response was followed with an increase in Go onset time (thus leaving less time to respond and
206 making the next trial more challenging; see detailed description in the methods section).

207 To identify unique response patterns, we performed an exploratory analysis of the RT
208 distribution during Go trials. Before analyzing RT in the CAT task, invalid trial trials were
209 excluded. Resulting in the exclusion of missed trials, in which participants failed to make a
210 response within the allocated 1500ms from stimulus onset timeframe (0.90% of trials), and trials
211 in which the response was shorter than a threshold of 100ms (0.02% of trials), as these trials are
212 likely to be indicative of inattentional response. In total, 1731 trial out of 188,224 Go trials were
213 excluded (0.92% of trials).

214 Examining the RT density distribution in the training task as a function of training run,
215 revealed a distinct pattern, wherein as training progressed, participants' RTs were less
216 homogeneous and started to form a growing peak of early RTs (Figure 2a). In all 29 experiments,
217 the Go cue changed in each trial, according to participants' performance (see methods), thus, a
218 more informative measurement of RT was the time from cue onset (RT minus cue-onset), referred
219 to here as *effective RT*. In earlier training runs, when the Go stimuli were associated for the first
220 time with the Go cue, most effective RTs were clustered around a unimodal center, approximately
221 300ms following the Go signal onset (effective RT M = 293ms) with 99% of effective RTs larger
222 than 145ms. We use here this empirical quantile of 145ms as threshold to define rapid *anticipatory*
223 *responses*, as these responses can be elicited when the participants predicted a go cue would
224 appear. As training progressed, a consistent pattern appeared in the effective RT data - the main
225 peak of the RT distribution was reduced, while a growing portion of participants' responses
226 consisted of faster anticipatory RTs, in many cases preceding the Go cue onset (Figure 2b). For
227 example, while the proportion of anticipatory responses comprised only 1% of RTs in the first
228 training run, they comprised 2.7% of RTs by the 5th run, 12.2% of the trials by the 10th run, 20.5%
229 of trials by the 15th run, and finally 29.4% of trials in the final 20th run (aggregated across all
230 samples).

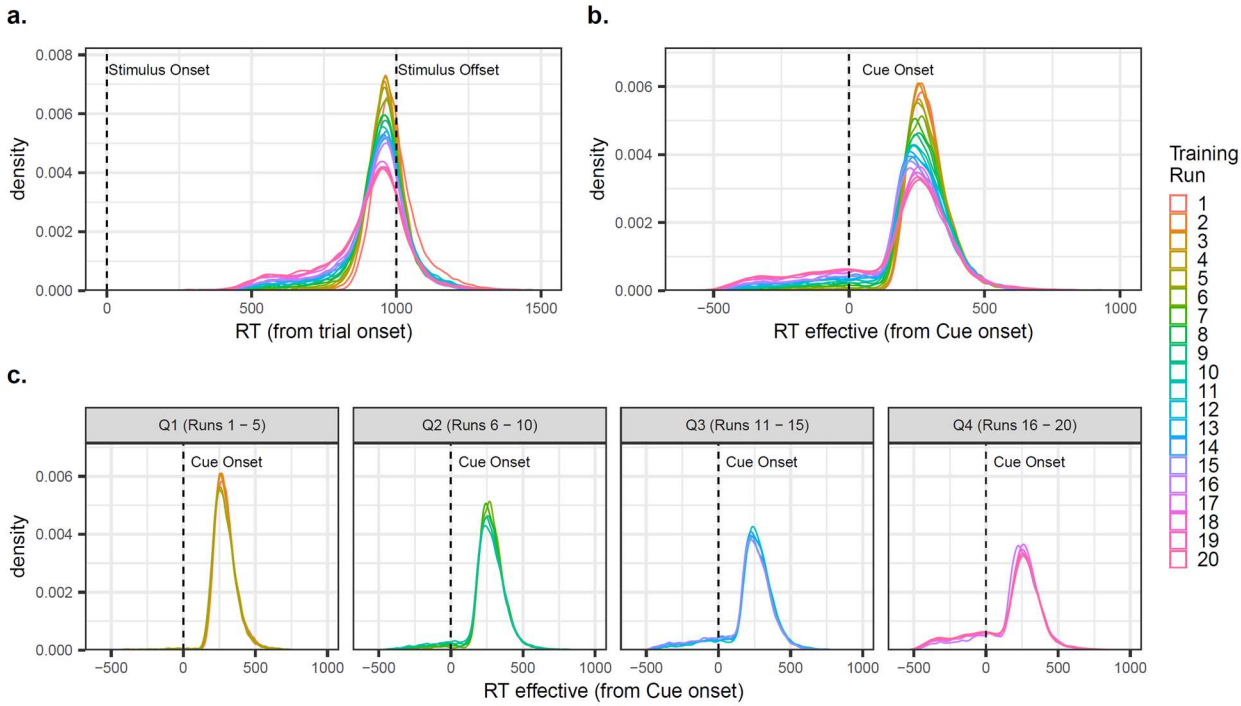


Figure 2. Reaction time distribution of data pooled from meta-analysis with 29 studies. Density plots of (a) RTs time-locked to the trial onset and (b) RTs time-locked to the varying Go signal onset (*effective RT*). Color indicates training run (repetition). As training progressed, the RT distribution shifted from a cue-dependent unimodal distribution to left-tailed distribution with increasing proportion of early anticipatory responses. (c) a clear differentiation in effective RT is apparent when splitting the data (from Figure 2b) to four quartiles, according to the training run (each quartile indicating five consecutive training runs).

Thus, RT data suggest a temporal-dependent response pattern. As training progress, participants tended to rely less on the Go cue (late cue-dependent RTs) and generated more early anticipatory responses. Interestingly, the mode of the RT distribution seemed to remain stable, and only the relative proportion of the two distributions changes. We hypothesized that this transition pattern from the cue-dependent response to a stimulus-triggered anticipatory response reflected a process of learning about the stimuli that could be associated with behavioral change in the later preference probe. Thus, we next sought to develop a model to characterize the strength of this effect within-individual.

Individualized-learning computational model. Based on the unique RT pattern observed in the exploratory analysis of CAT task data, we developed a novel computational Bayesian model. To capture the form of RT distribution, we modeled the effective RT as a mixture of two Gaussian distributions, with two different means (μ_1 and μ_2 free parameters), two standard deviations (σ_{ε_1} and σ_{ε_2}), and a mixture proportion ($\Theta_{i,t}$) indicating the relative mixture proportion for participant$_i$

at $trial_t$. The model was implemented with the Stan probabilistic programming language [38], which optimized the model parameter estimates using a Markov-chain Monte Carlo (MCMC) approach. See methods section for a detailed description of the model and Supplementary Code 1. The first Gaussian was restricted to have lower mean than the second Gaussian to maintain their correspondence across different MCMC sampling chains (i.e., $\mu_1, \sigma_{\varepsilon_1}$ would correspond with the early anticipatory RT distribution). To capture the gradual change in anticipatory responses over training, the mixture proportion was modeled as a time-dependent variable could be expressed by the following formula:

$$\Theta_{i,t} = \phi\left(\theta_0 + \theta_{slope_i} Run_{i,t}\right)$$

Where the mixture probability $\Theta_{i,t}$ for $participant_i$ at $trial_t$ was modeled as a linear function (with a fixed θ_0 intercept and random participant-level θ_{slope_i} free paramter) of training run (where the $Run_{i,t}$ ranged between [0,1]). The linear trend was scaled non-linearly to probability-appropriate values of [0, 1] range using a normal distribution cumulative distribution function (normal CDF; denoted with ϕ), resulting in a sigmoid-like link function. See further details in the methods section.

For easy interpretation of the parameters, the θ_0 intercept term was fixed to -3.1, corresponding with $\Theta_{i,t} = \phi(-3.1) \approx 0.1\%$ mixture proportion at the very first run for all participants. Thus, higher θ_{slope_i} parameter estimate indicates an individual participant with faster transition from slow cue-dependent RTs to fast anticipatory responses, which corresponds with larger $\Theta_{i,t}$ mixture proportion at the very last run. E.g., parameter estimate of $\theta_{slop\ \ i}$= 3.1, indicated that at the last run ($Run_{i,t}$ = 1), the mixture proportion of anticipatory responses would be $\phi(0) = 50\%$, while $\theta_{slop\ \ i}$= 4.745 would correspond with mixture proportion of $\phi(1.645) \approx$ 95%, at the last run. We hypothesized that this individualized parameter would be indicative of improved learning in the training phase of CAT and would also be positively associated with stronger preference modification effect, measured in the subsequent probe phase.

After running four independent chains, the Bayesian model converged to a stable solution (see Supplementary Figure 1). The converged model estimated effective RTs as a time-dependent mixture of two RT distributions: one of early anticipatory responses (μ_1 = 107.50ms, 95%CI [103.77, 111.21], σ_{ε_1} = 276.30ms, 95%CI [274.13, 278.42]) and late cue-dependent responses (μ_2 = 286.12ms, 95%CI [285.67, 286.58], σ_{ε_2} = 72.61ms, 95%CI [72.23, 73.00]). Overall, participants

demonstrated an increase in proportion of anticipatory responses as training progressed, as manifested in the positive group-level parameter (θ_{slope} = 4.19, 95%CI [3.84, 4.54]), with variation between participants ($\sigma_{\theta_{slope}}$ = 5.14, 95% CI [4.85, 5.46]), indicating some participants made faster transitions and some generated nearly only cue-dependent responses (non-positive θ_{slope_i} estimates).

Posterior predictive checks (simulated distributions of RT based on estimated model parameters) revealed a good fit of the model to the actual data. Simulated posterior distributions recreated the patterns observed empirically, showing more rapid transition to anticipatory responses in participants with higher $\theta_{slop\ \ i}$ parameter estimate (Figure 3; Supplementary Figure 2).

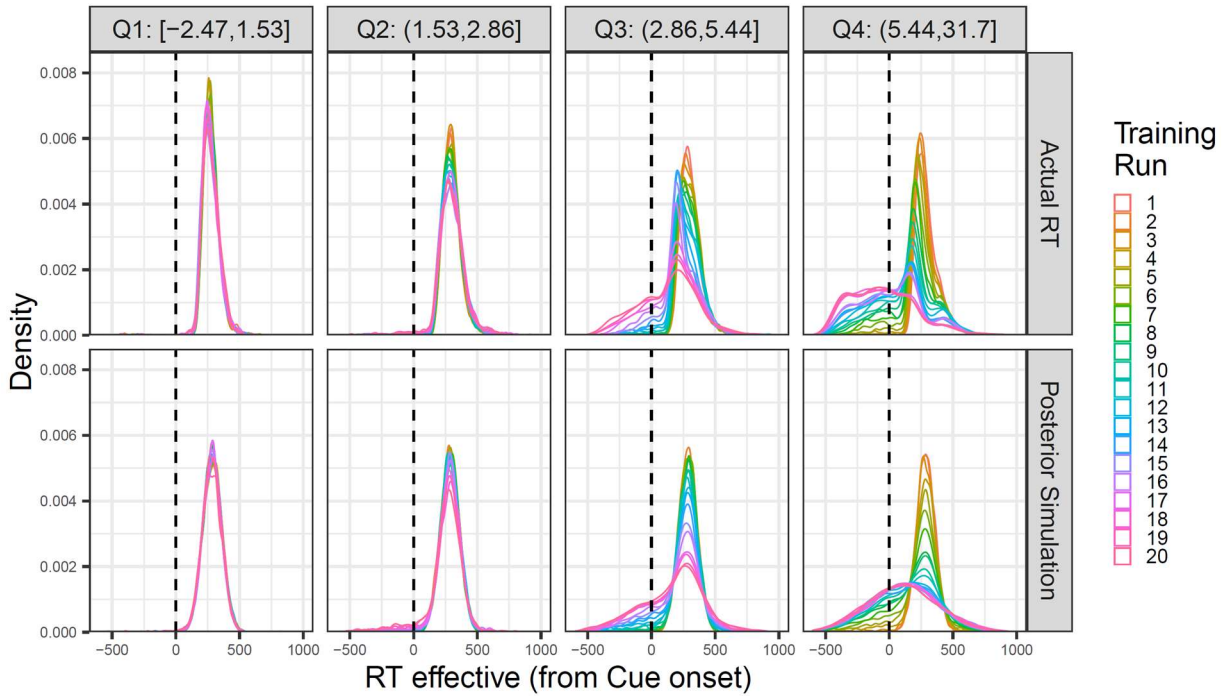


Figure 3. Actual RT distributions versus simulated posterior distributions, by θ_{slope_i} quantile group. Participants were categorized into four equal quantile groups, according to their parameter estimates (denoted here as Q1-Q4; columns). Participants with higher parameter estimates were characterized with faster transition to anticipatory responses (top row). Posterior simulated RT distributions using mixture of Gaussians (bottom row) recreated relatively well this transition pattern. Vertical dashed line represents cue-onset. See also Supplementary Figure 2 for more detailed comparison.

Learning parameter association with choices. Previous findings with CAT consistently showed that preferences for Go stimuli were enhanced following CAT, as manifested in choice

behavior during the probe phase. When presented with a choice between a Go stimulus and a NoGo stimulus of similar initial value, participants consistently chose the Go stimulus [12,14–21]. To test the hypothesis that this probe performance is associated with RT patterns during CAT, we examined whether variation in the slope parameter fit to RTs correlated with probe performance. Note that, under the model, the slope parameter controls the final proportion of anticipatory responses, which we understood as a measure of the extent of learning about the stimulus achieved by the endpoint of CAT. To evaluate the association of the θ_{slope_i} parameter with preference-change effect following CAT, we analyzed the proportion of probe trials in which participants chose the Go over the NoGo stimulus, as a per-participant linear function of θ_{slope_i} (mixed logistic regression model, including random intercept and slope terms for the 29 experiments; see methods).

The meta-analysis showed θ_{slope_i} was positively associated with the preference modification effect – i.e., participants with higher θ_{slope_i} parameter estimate, also demonstrated greater odds of choosing Go stimuli (OR = 1.04, 95% CI = [1.02, 1.06], $Z = 5.17$, $p = 2.3E^{-7}$; one-sided mixed model logistic regression; Figure 4). The model's intercept was significantly greater than zero, i.e., even when extrapolated to very low θ_{slope_i} value, the model forecasted enhanced preference for Go stimuli (intercept odds = 1.26, 95% CI = [1.18, 1.34], $Z = 7.03$, $p = 2.0E^{-12}$).

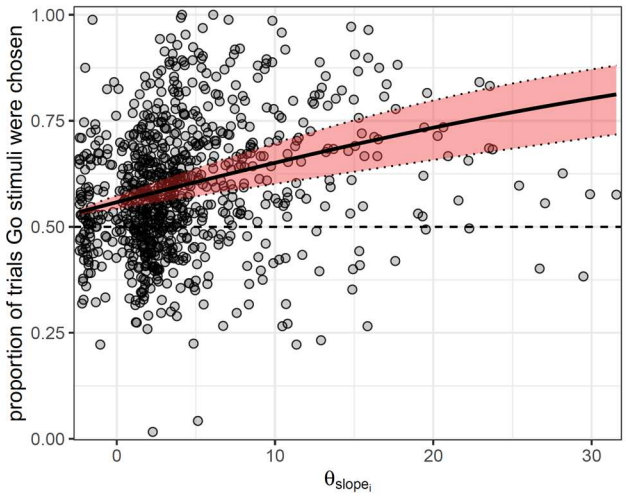


Figure 4. Meta-analysis results - computational marker and preference modification effect. Participants who transitioned faster to anticipatory response during cue-approach training (larger $\theta_{slop\ \ i}$ estimates) also demonstrated stronger preference modification effect (proportion of trials Go stimuli were chosen). Trend line and surrounding red margins represent estimated preference modification effect and 95% CI, respectively (mixed model logistic regression). Dots represent individual participants.

The estimated variances associated with the fixed-effect terms, random-effect terms and residuals were used to evaluate two R^2 scores for the generalized (logistic) linear mixed mode: a marginal $R^2_{GLMM(m)}$ (representing the relative proportion of variance associated with the fixed-effects) and a conditional $R^2_{GLMM(c)}$ score (representing the relative proportion of variance associated with the fixed-effects and random-effects; see more details in the methods section). The θ_{slope_i} fixed effect accounted for $R^2_{GLMM(m)}$ = 0.311 of the variance. With the random intercept and slopes, the fixed and random effects combined accounted for $R^2_{GLMM(c)}$ = 0.828 of the overall variance.

Stimulus-specific learning parameter. While the previous model aggregated the data within participants, in fact, each participant has learned the Go cue association with several different Go stimuli. Thus, it is possible that learning was not uniform across the entire training stimuli, meaning every participant had a slightly different learning rate for each of the Go stimuli she or he encountered. In an additional model, we aimed to expend our model by examining behavior on a more granular scale – focusing on the variability in RT and choice for individual Go stimuli within participant. Accordingly, the computational model was elaborated by decomposing the per-subject learning parameter $\theta_{slop\ \ i}$ into a set of subject- and stimulus-specific parameters $\theta_{slop\ \ i,s}$, indicating the transition speed to anticipatory RTs of each Go stimulus$_s$ encountered by participant$_i$ during training. The model was otherwise identical in design (see methods). This reanalysis of the data was conducted after the collection and analysis of the following two studies which are reported in the current work.

The reanalyzed model converged around slightly different parameter estimates: estimated early anticipatory RT distribution was characterized with lower mean (μ_1 = 88.30ms, 95% CI [84.1, 92.34], σ_{ε_1} = 278.2ms, 95% CI [275.84, 280.65]), late cue-dependent responses were similar (μ_2 = 289.31ms, 95% CI [288.82, 289.79], σ_{ε_2} = 75.64ms, 95% CI [75.23, 76.05]), and the $\theta_{slop\ \ i,s}$ parameters tended to be lower but with greater variability (θ_{slope} = 2.18, 95% CI [2.01, 2.34], $\sigma_{\theta_{slope}}$ = 5.95, 95% CI [5.76, 6.16]; see Supplementary Figure 3).

Similarly to the previous logistic regression analysis with per-participant parameter, using the current analysis' per-stimulus parameters also significantly predicted choices of the corresponding stimuli in the subsequent probe phase (Intercept: Odds = 1.45, 95%CI [1.36, 1.54], p = 7.28E^{-30}; $\theta_{slope_{i,s}}$ OR = 1.02, 95%CI [1.01, 1.03], Z = 3.65, p = 1.3E^{-4}; one-sided mixed model

logistic regression). Thus, Go stimuli which were characterize with larger proportion of anticipatory responses were also chosen more during the probe phase. The models' fixed effects accounted for $R^2_{GLMM(m)}$ = 0.011 of the total variance and the fixed effect with mixed effects accounted for $R^2_{GLMM(c)}$ = 0.775 of the total variance.

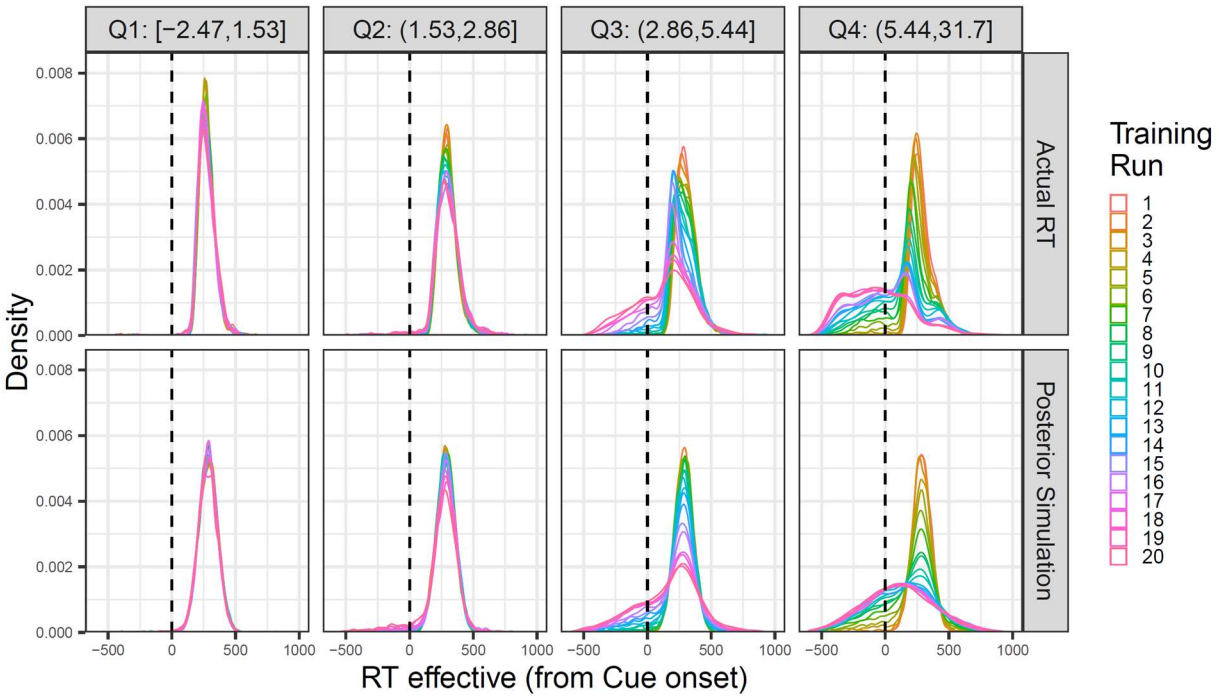
However, this overall effect comprises both between-participant and within-subject, between-stimulus contributions. To examine the unique contribution of $\theta_{slope_{i,s}}$ above and beyond participant-level $\theta_{slop\ \ i}$ parameter (which was evaluated in the previous analysis), in a third logistic regression model we included both per-participant and per-stimulus parameters from the two different Stan models, as independent variable (and their random effects; see methods) to predict choices during probe. Our results indicate that each parameter provided some unique contribution above and beyond the other parameter (Intercept: Odds = 1.37, 95%CI [1.27, 1.47], p = 2.38E^{-18}; stimulus-level $\theta_{slope_{i,s}}$: OR = 1.02, 95%CI [1.01, 1.03], Z = 3.34, p = 4.2E^{-4}; participant-level θ_{slope_i}: OR = 1.01, 95%CI [1.00, 1.03], Z = 2.07, p = 0.019). Thus, stronger learning of anticipatory RTs, both per-individual and per-stimulus within individual, were associated with increased odds of choosing Go stimuli.

Interim discussion

Examining RT patterns during CAT revealed for the first time a distinct time-dependent RT pattern, in which participants gradually transition from slow cue-dependent responses to rapid anticipatory RTs. Using Bayesian modeling, we were able to quantify this RT transition pattern with stable parameter estimates, of which the θ_{slope_i} parameter provided a promising computational marker to evaluated individualized differences in learning during training. Furthermore, as expected, this computational marker was also found to be associated with the preference modification effect in the subsequent probe phase, as participants with a more robust learning marker also demonstrated stronger preference modification effect. Examining the data on a stimulus-level learning parameter, demonstrated an improved explanatory power for the variability in probe choices, above and beyond the wider participant-level parameter. Since the computational marker was calculate based on measurements that preceded the probe phase, it holds a potential to provide predictive marker for future preference modification.

Several challenges were noted in modeling the original experimental data with the new computational model, largely owing to the fact that the original experimental procedures were not designed with the present analyses in mind. First, in the training task, Go cue onsets were derived

based on performance using a staircase procedure. Participants who performed well and responded rapidly were presented in the next trials with a more challenging cue, appearing later into the trial, while participants who performed poorly were exposed to earlier occurring (less difficult) cues. Consequently, anticipatory responses were not time-locked to the same event as the late cue-dependent responses, providing a challenge for a model trying to differentiate between the two. For example, participants with very poor performance could have been presented with a cue at 500ms from stimulus onset and respond at 750 from stimulus onset, while a well-performing participant could encounter a challenging 800ms cue onset delay but respond before cue-onset at 750ms from stimulus onset, making on the surface the two response patterns hard to distinguish and introducing a potentially confounding source of variation across participants. This challenging effect may have led to some counter-intuitive results; for instance, due to high variability in anticipatory responses, the fitted model predicted that very late responses (RT effective > 500) would be more likely in the fastest learners' late runs, compared with earlier runs or with slower learners (



).

Some lack of fit of the predicted data could also be indicative that the assumption that anticipatory RTs are normally distributed around the same mean for all participants is unlikely. Nonetheless, the large heterogenous data structure (with over 800 participants from 29 different experiments) provided a computational challenge when trying to fit more complex models (e.g., with random effects per participant on this parameter). The current model was selected as a reasonable compromise that provided sensible results.

Furthermore, the task instructions of CAT specifically mentioned that participants are required to respond *after* cue-onset. An additional crucial factor which could not be controlled in

the current study was how rigorously participants complied with this instruction. It is possible that some participants might have learned the stimulus-cue association well and could have produced anticipatory responses but were reluctant to deviate from the task's guidelines. If so, their θ_{slope_i} would not reliably reflect their actual learning of the task. It is important to note that although the logistic regression results showed a significant positive association of choices with participant-level even stimulus-level individualized learning parameters, significant preference change remains even when the model extrapolated for no learning effect, and the effect sizes reported above are quite modest. Thus, suggesting of additional contributions to the choice pattern which are not otherwise accounted for by the computational marker. The small effect sizes might be due to the inaccurate measurement of additional constructs within the computation parameter, such as the instruction adherence and interaction with performance-based dynamic cue onset, which were proposed here.

Despite these drawbacks, our unique model provided a prospective computational marker for learning and subsequent preference changes. Based on the promising results of Study 1 meta-analysis and considering the challenges raised above in fitting the computational model to the CAT task data, in Study 2, we devised a novel design of the CAT procedure, which was tested with two independent experiments, a smaller preliminary study (n = 25) and a larger pre-registered replication study (n = 59). In these new experiments we addressed the limitations of the previous study design. We also modified the training design to answer a more directional hypothesis which does not only examine correlational association, but also try to provide evidence for more consequential directionality, by inducing a differential behavioral change.

Study 2: Novel CAT design – preliminary and replication experiments

A novel experimental design was tested in two pre-registered experiments - a preliminary experiment and a larger replication experiment. Based on the conclusions from Study 1, the CAT procedure and instructions were altered to optimize the task for the Bayesian computational framework and manipulate behavior in accordance with the hypothesized cognitive mechanism.

In the training task, the Go cue onset time was fixed at 850ms from stimulus onset and participants were clearly instructed that they were permitted to make anticipatory responses before cue onset. Furthermore, based on the conclusions from the meta-analysis study, we introduced within the training task a manipulation of the predictability of the stimulus-Go cue contingency, which we expected would affect the θ_{slope_i} parameter and subsequently also manipulate the

preference modification effect. Of the Go stimuli, half of the stimuli were always associated with the Go cue (100% contingency condition), while the rest of the Go stimuli were only associated in half of the presentations with the Go cue (50% contingency condition). The rest of the stimuli were never associated with a Go cue (NoGo stimuli; See methods section for detailed description of the new design).

We hypothesized that in the 50% contingency condition, learning the association of Go stimuli with the Go cue would be more challenging, and would therefore be identified by a lower θ_{slope_i} learning parameter. We also hypothesized that manipulating learning efficacy during training would therefore induce a differential preference modification effect in the subsequent probe phase, with more robust preference modification for Go stimuli trained in the 100% contingency condition, compared with the more challenging 50% contingency condition.

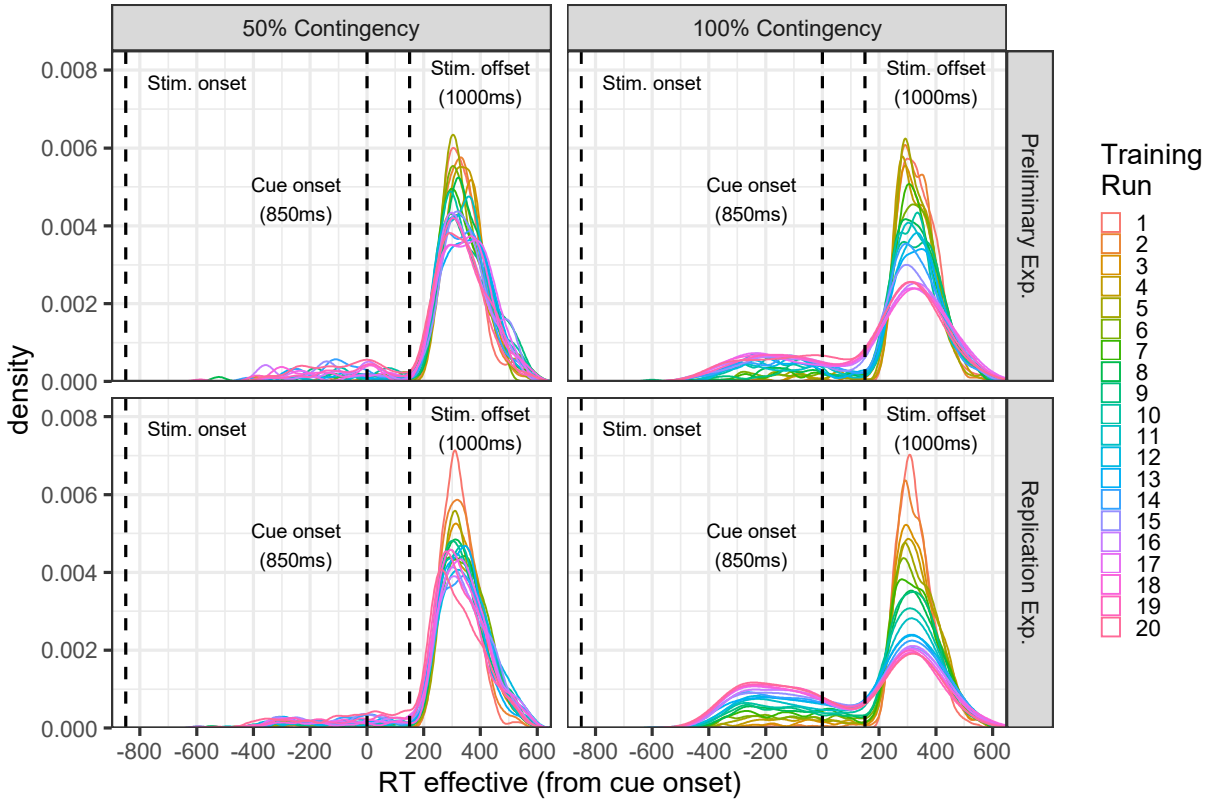
The new task design was first tested in a preliminary (not preregistered) study with $n = 20$ valid participants, aimed to evaluate the efficacy of the new task and the replicability of the meta-analysis conclusions. Following the preliminary experiment, we run an additional pre-registered direct replication experiment with a larger sample size ($n = 59$), a size chosen based on the preliminary experiment results. As the two experiments were identical in design, they are reported together in all sections.

Results

Reaction times in the training task. As in the meta-analysis study, RT patterns of the training task were examined. Before examining RT patterns, we removed Go trials where participants did not respond at all (preliminary exp. = 6.33%. replication exp. 5.97% of Go trials) or trials with an unlikely fast RT (RT< 250ms, preliminary exp. = 0.31%, replication experiment 0.21% of trials). In total, 636 (of 9600; 6.62%) and 1747 (of 28,320; 6.17%) of trials were excluded from the preliminary and replication experiments, respectively.

In both experiments, we were able to replicate the RT pattern observed in the meta-analysis study – while in early training runs, participants mostly relied on cue-dependent RTs, as training progressed, an increased portion of RTs were of earlier anticipatory RTs (Figure 5). Examining mean (*SD*) effective RT at Run 1, revealed similar RTs in all contingency conditions - preliminary exp.: $M_{50\%Cont.}$ = 326.88ms (85.13ms), $M_{100\%Cont.}$ = 336.60ms (64.39ms); replication exp.: $M_{50\%Cont.}$ = 322.49ms (63.68ms), $M_{100\%Cont.}$ = 319.90ms (78.88ms). As training progressed,

475 participants were able to anticipate the cue and respond before cue-onset. Examining the
476 differential pattern between the 100% contingency condition and the 50% contingency condition
477 showed, as expected, that participants initiated more anticipatory responses in the 100%
478 contingency condition, compared with the 50% contingency condition, resulting in faster mean-
479 RTs at Run 20 - preliminary exp.: $M_{50\%Cont.}$ = 253.27ms (187.65ms), $M_{100\%Cont.}$ = 175.55ms
480 (245.5ms); replication exp.: $M_{50\%Cont.}$ = 256.57ms (188.94ms), $M_{100\%Cont.}$ = 81.39ms
481 (271.99ms); see Figure 5.



482
483 Figure 5. Density plots of effective RT distribution (time-locked to cue onset) in Study 2
484 (preliminary experiment on top row and replication experiment on bottom row). The Go cue onset
485 was fixed at 850ms from stimulus onset - vertical dashed lines represents the trial onset (time 0),
486 cue-onset, and trial offset (850ms and 1000ms from trial onset, respectively). As training
487 progressed (indicated by the training run in color), responses were less homogeneous following
488 cue onset, with increased proportion of anticipatory responses. The transition to anticipatory
489 responses was more robust in the 100% contingency condition (right column) compared with the
490 50% contingency condition (left column).
491
492 ***Computational marker of learning.*** As in the meta-analysis study, RTs within each
493 condition were modeled with a time-dependent mixture model, implemented with Stan Bayesian
494 framework. RT data within each condition (100% contingency and 50% contingency) were fitted

using two Gaussian distribution parameters (two means and two SDs). A θ_{slope_i} time dependent parameter was modeled for each condition and determined the rate of transition from cue-dependent responses to anticipatory responses of individual participants. Since the Go cue in Study 2 was fixed at 850ms, the effective RT (relative to stimulus rather than cue) measurement perfectly correlated with actual RT; thus, we used only the actual RT in the model.

In both the preliminary and replication experiment, the computational model converged around similar hyperparameter estimates (see MCMC trace plot and posterior simulations fit in Supplementary Figure 5 and Supplementary Figure 6, respectively). Cue-dependent responses were characterized with later mean RT and smaller variance, compared with earlier anticipatory responses (preliminary experiment: μ_1 = 735.07, 95%CI [677.14, 792.87], σ_{ε_1} = 140.14, 95%CI [130.13, 153.49], μ_2 = 1190.23, 95%CI [1188.46, 1192.03], σ_{ε_2} = 78.11, 95%CI [76.78, 79.47]; replication experiment: μ_1 = 740.44, 95%CI [710.10, 771.76], σ_{ε_1} = 140.35, 95%CI [135.79, 144.74], μ_2 = 1187.46, 95%CI [1186.25, 1188.68], σ_{ε_2} = 80.30, 95%CI [79.46, 81.15]). As the cue onset was fixed in Study 2 at 850ms, the mean cue-dependent RTs of approximately 1190ms corresponded with an effective RTs center of approximately 340ms in both experiments. Anticipatory RT mean varied considerably between participants, as identified by the σ_{μ_1} parameter, which modeled between-participant variability in μ_{1_i} parameter estimates (preliminary experiment: σ_{μ_1} = 110.81, 95%CI [73.29, 166.69]; replication experiment: σ_{μ_1} = 110.99, 95%CI [90.92, 136.34]).

Transition from cue-dependent to anticipatory responses, as captured by θ_{slope_i} parameter estimates of the preliminary experiment, indicated faster transition in the 100% contingency condition, compared with the 50% contingency condition ($\theta_{slope100\%}$ = 1.95, 95%CI [1.03, 2.75], $\theta_{slope50\%}$ = 0.90, 95%CI [-0.01, 1.68]; mean difference in θ_{slopes} = 1.04, 95%CI = [-0.17, 2.23]), with prominent variability between participants ($\sigma_{\theta_{slope100\%}}$ = 1.98, 95%CI [1.45, 2.67], $\sigma_{\theta_{slope50\%}}$ = 1.79, 95%CI [1.21, 2.57]). This effect was replicated even more distinctly in the larger replication experiment ($\theta_{slope100\%}$ = 3.25, 95%CI [2.54, 3.89], $\theta_{slope50\%}$ = 1.54, 95%CI [1.11, 1.92]; mean difference in θ_{slopes} = 1.70, 95%CI = [0.90, 2.47]; $\sigma_{\theta_{slope100\%}}$ = 2.82, 95%CI [2.37, 3.42], $\sigma_{\theta_{slope50\%}}$ = 1.45, 95%CI [1.16, 1.82]). The 95% credible interval of the difference between

the two conditions $\theta_{slop\ i}$ parameters indicated a general trend in the preliminary experiment which was distinct in the replication experiment.

Preference modification in probe task. Based on the results of the meta-analysis in Study 1, we hypothesized that manipulating the Go cue contingency would correspondingly manipulate the preference modification effect, manifested as more pronounced preference modification for stimuli in the 100% contingency condition, compared with the 50% contingency condition.

As expected, in the preliminary experiment, participants chose Go stimuli over NoGo stimuli above chance level, both in the 50% contingency condition (prop. = 57.11%, $Z = 2.85$, $p = 0.002$, odds = 1.33, 95% CI [1.09, 1.62]; one-sided logistic mixed model), as well as in the 100% contingency condition (prop. = 64.45%, $Z = 3.40$, $p = 3.4E^{-4}$, odds = 1.81, 95% CI [1.29, 2.56]). The preference modification effect was more robust in the 100% contingency condition, compared with the 50% contingency condition ($Z = 1.92$, $p = 0.028$, OR = 1.36, 95% CI = [0.99, 1.87]). These effects were prominently replicated in the larger replication experiment (50% contingency condition: prop. = 58.41%, $Z = 5.50$, $p = 1.9E^{-8}$, odds = 1.40, 95% CI [1.24, 1.59]; 100% contingency condition: prop. = 66.31%, $Z = 6.41$, $p = 7.1E^{-11}$, odds = 1.97, 95% CI [1.60, 2.42]; condition difference-effect: $Z = 3.81$, $p = 7.0E^{-5}$, odds = 1.40, 95% CI [1.18, 1.67]; one-sided logistic mixed model), see Figure 6.

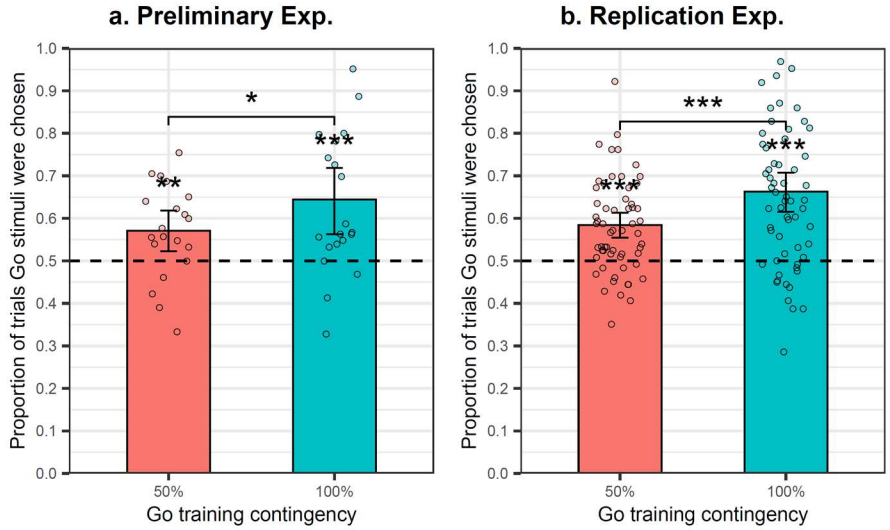


Figure 6. Probe results. Proportion of trials participants chose Go stimuli over NoGo stimuli of similar initial value, in the preliminary experiment (a) and replication experiment (b). Dots represent individual participants, error-bars represents 95% CI based on a mixed model logistic regression. Participants demonstrated enhance preference both for Go stimuli in the 50% contingency condition and to a larger extent in the 100% condition. Statistical significance is

denoted with asterisks (* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$; one-sided mixed model logistic regression). Dashed line represents 50% chance level.

Prediction of probe using computational model. Most importantly, we aimed to examine whether the preference modification effect could be predicted using the individually fitted θ_{slope_i} parameter estimated for each participant. Adding the individual learning parameter as an additional independent variable to the logistic regression model revealed that indeed θ_{slope_i} parameter estimated from the training task, could be used to predict preference modification in the subsequent probe task (Figure 7). In the preliminary experiment a significant contribution was found for θ_{slope_i} in the 50% contingency condition (log-OR = 0.12, $Z = 2.30$, $p = 0.011$, OR = 1.13, 95% CI [1.02, 1.26]; one-sided mixed model logistic regression), as well as in the 100% contingency condition (log-OR = 0.26, $Z = 3.59$, $p = 1.7E^{-4}$, OR = 1.29, 95% CI [1.12, 1.49]). No significant difference was found between the slopes under the two conditions (log-OR = 0.13, $Z = 1.52$, $p = 0.13$, OR = 1.14, 95% CI [0.96, 1.36]; two-sided mixed model logistic regression). These results were replicated also in the larger replication experiment, where θ_{slope_i} parameter estimates predicted the preference modification effect, both in the 50% contingency condition choices (log-OR = 0.10, $Z = 2.39$, $p = 0.008$, OR = 1.11, 95% CI [1.02, 1.20]; one-sided mixed model logistic regression), as well as in the 100% contingency condition (log-OR = 0.11, $Z = 3.49$, $p = 2.4E^{-4}$, OR = 1.12, 95% CI [1.05, 1.19]). No differential slope effect was found between the two conditions (log-OR = 0.01, $Z = 0.26$, $p = 0.79$, OR = 1.01, 95% CI [0.93, 1.10]; two-sided mixed model logistic regression).

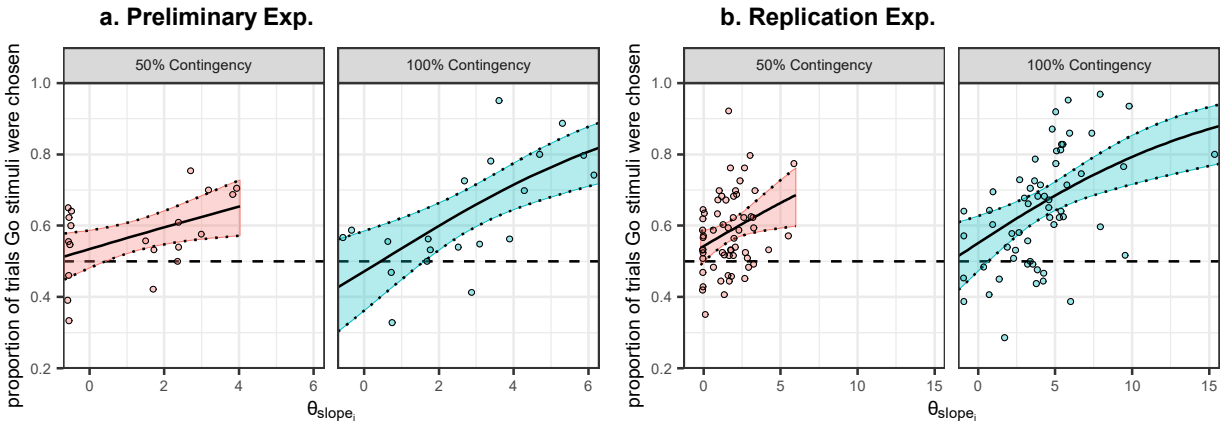


Figure 7. Association of probe choices with training computational marker in study 2. Larger $\boldsymbol{\theta_{slope_i}}$ estimates parameter estimates were positively associated with subsequent preference modification in probe phase, both in the preliminary experiment (a) and in the larger replication experiment (b). The 100% contingency condition was characterized with larger $\boldsymbol{\theta_{slope_i}}$ values, and respectively also stronger preference modification effect. Trend line and surrounding color margins represent estimated preference modification effect and 95% CI, respectively (mixed model logistic regression). Dots represent individual participants. Horizontal dashed line represents 50% chance level.

Examining the sizes of estimated fixed-effect, random-effect and residual variances, revealed that the model accounted for a large portion of the variance in participants' choice pattern in both experiments, both when examining the contribution of the fixed effects only (preliminary exp.: $R^2_{GLMM(m)}$ = 0.421; replication exp.: $R^2_{GLMM(m)}$ = 0.223), and furthermore when examining the joint contribution of fixed and random effects (preliminary exp.: $R^2_{GLMM(c)}$ = 0.869; replication exp.: $R^2_{GLMM(c)}$ = 0.872).

To examine the overall effect of contingency condition above and beyond θ_{slope_i}, using a likelihood ratio test, the logistic regression model was compared with a restricted (nested) model that did not contain contingency fixed effects (contingency intercept and slope interaction; restriction of 2 parameters). In both experiments, comparing the two models revealed no significant contribution of contingency above and beyond θ_{slope_i} (preliminary experiment: $\Delta AIC_{(rest.-full)}$ = -1.64, $\chi^2_{(2)}$ = 2.36, p = 0.307; replication experiment: $\Delta AIC_{(rest.-full)}$ = -3.61, $\chi^2_{(2)}$ = 0.39, p = 0.824; two-sided likelihood ratio test).

Stimulus-specific learning parameter. In an additional analysis (mentioned but not detailed in the preregistration), we extended this analysis to stimulus-level effects. Each Go stimulus$_s$ presented to participant$_i$ was fitted a $\theta_{slope_{i,s}}$ parameter, which was expected to capture

within-participant variability in learning - i.e., model for difference in learning of stimuli within a contingency condition. We aimed to use these participant and stimulus-specific learning parameters as independent variables in a mixed model logistic regression (with a random intercept, random slope for contingency and random slope for $\theta_{slope_{i,s}}$ independent variables, see methods for full description of the statistical models).

Fitting a full Stan model for both stimuli of 50% and 100% contingency simultaneously did not converge. We hypothesized that the full model, with 2 conditions × 16 stimuli × n participants inter-dependent $\theta_{slope_{i,s}}$ parameters was too complex to resolve using MCMC process. Thus, we decided to reduce complexity by eliminating the participant-level effects on the mean RT for the early anticipatory responses (estimating only group-level μ_1, without participant-level μ_{1_i} parameters; as was done in the previous meta-analysis study). We further split the data based on contingency condition and analyzing each contingency as an independent dataset. Using this approach resolved the convergence issues. This solution for an unexpected issue was not preregistered or planned prior to data analysis.

Examining the association between the $\theta_{slope_{i,s}}$ parameter estimates and the participant-level $\theta_{slop\ \ i}$ parameter estimates of the previous model, revealed high similarity (preliminary exp.: r = 0.84; replication exp.: r = 0.80). In addition, participants with negative θ_{slope_i} parameter estimates had very small variance (SD < 0.1) in their $\theta_{slope_{i,s}}$ estimates (see Supplementary Figure 7). When such $\theta_{slope_{i,s}}$ were introduced in the mixed-model logistic regression, the low within-participant variability caused convergence warnings. Therefore, all participants with θ_{slope_i}< 0.2 (which is equivalent to SD < 0.1 of $\theta_{slope_{i,s}}$) in either contingency condition, were excluded from the analysis. This resulted in the exclusion of nine participants in the preliminary experiment and 14 participants in the replication experiment.

Analyzing the remaining 11 and 45 participants in the preliminary and replication experiment, resulted in similar conclusions, as found with the participant-level θ_{slope_i} parameter estimate. Overall, in both contingency conditions, a significant positive association was found between $\theta_{slope_{i,s}}$ and preference of Go stimuli over NoGo stimuli (preliminary experiment - 50% contingency: Z = 4.34, p = 7.0E^{-6}, OR = 1.33, 95% CI [1.17, 1.52]; 100% contingency: Z = 2.85, p = 0.002, OR = 1.21, 95% CI [1.06, 1.38], overall model $R^2_{GLMM(m)}$ = 0.269, $R^2_{GLMM(c)}$ = 0.631;

replication experiment - 50% contingency: $Z = 3.14$, $p = 8.4E^{-4}$, OR = 1.09, 95% CI [1.03, 1.15]; 100% contingency: $Z = 5.09$, $p = 1.7E^{-7}$, OR = 1.09, 95% CI [1.05, 1.12], overall model $R^2_{GLMM(m)}$ = 0.147, $R^2_{GLMM(c)}$ = 0.475; one-sided mixed model logistic regression; Figure 8). No significant differences were found between the slopes of the two conditions (preliminary experiment: $Z = -1.35$, $p = 0.18$, OR = 0.91, 95% CI [0.79, 1.04]; replication experiment: $Z = -0.08$, $p = 0.94$, OR = 1.00, 95% CI [0.95, 1.05]; two-sided mixed model logistic regression).

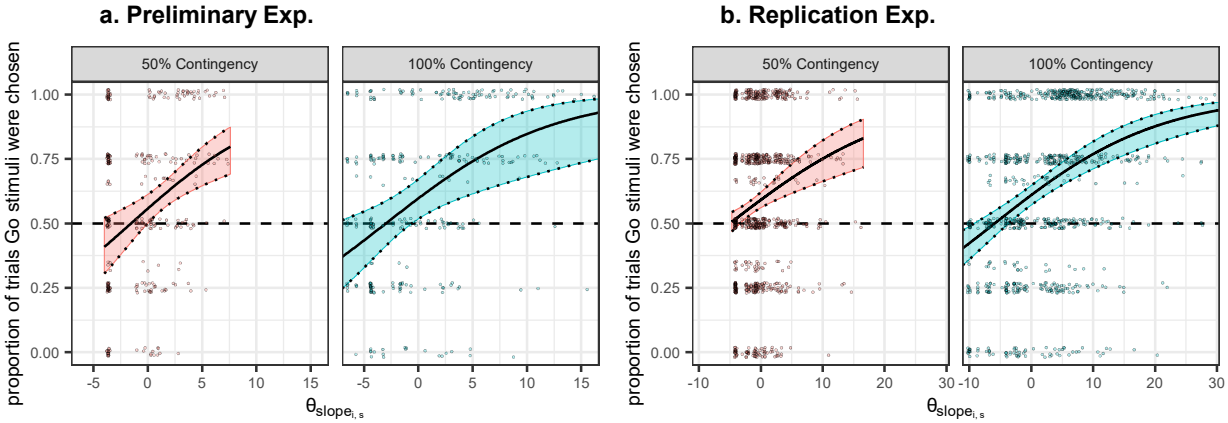


Figure 8. Association of probe choices with stimulus-level computational marker in study 2. Larger $\theta_{slope_{i,s}}$ estimates parameter estimates were positively associated with subsequent preference modification in probe phase, both in the preliminary experiment (a) and in the larger replication experiment (b). Dots represent individual stimuli (small vertical jitter was added to better visualize high density areas). Trend lines and surrounding color margins represent estimated preference modification effect and 95% CI, respectively (mixed model logistic regression). Horizontal dashed line represents 50% chance level.

The contribution of stimulus-level $\theta_{slope_{i,s}}$ parameter to predicting preferences was examined by comparing a full model, containing both stimulus-level $\theta_{slope_{i,s}}$ and participant-level θ_{slope_i} individualized learning parameters to a restricted model containing only $\theta_{slo\ \ _i}$ as independent variable. The addition of $\theta_{slope_{i,s}}$ had a significant contribution both in the preliminary experiment (ΔAIC = 59.65, $\chi^2_{(9)}$ = 77.65, $p = 4.7E^{-13}$; likelihood ratio test), as well as in the replication experiment (ΔAIC = 93.97, $\chi^2_{(9)}$ = 111.97, $p = 5.8E^{-20}$).

Interim discussion

Based on the conclusions of the meta-analysis study, a novel preference modification procedure was designed and tested in Study 2. In the new design, two cue contingency conditions (50% and 100% contingency) were tested within-participant, which we hypothesize would

manipulate the difficulty of learning the stimulus-cue association during the training phase of the CAT paradigm. The introduction of the two contingency conditions allowed us to experimentally impact the $\theta_{slop\ \ i}$ individualized computational marker of learning, which in turn provided a reliable predictor for the subsequent preference modification effect in the probe phase. As expected, we found more a robust preference modification effect for Go stimuli in the 100% contingency condition, compared with the 50% contingency condition, as tested within-participants. The fact that a directed experimental intervention in the preceding training phase induced a differential effect in the probe phase, provides support for a directional causal impact of training on behavioral change. In light of these results, it is plausible to conclude that the association demonstrated in Study 1 between the individualized learning parameter and behavioral change might represent a causal relationship, in which improved learning in the training phase induces stronger behavioral change in the subsequent probe phase.

Furthermore, between-participant variability in the $\theta_{slo\ \ \ i}$ computational marker was also predictive of between-participant difference in preference modification, as measured during the subsequent probe phase. Participants with more robust θ_{slope_i} also demonstrated stronger preference for Go stimuli over NoGo stimuli of similar initial value. This effect was observed both in the 100% and 50% contingency conditions, with no significant difference in the effect between the two conditions. This suggests that θ_{slope_i} had a similar predictive pattern for both conditions, one unit increase in θ_{slope_i} resulted in similar increase in the odds of choosing Go over NoGo, above and beyond the contingency condition. However, as the 100% contingency condition was characterized with larger θ_{slope_i} parameter estimates, so was the probe phase characterized with greater odds of choosing Go stimuli (i.e., stronger preference modification effect). Taken together, these results provide evidence for novel means to both predict individual differences in learning and preference modification between participants, as well as means to manipulate this preference modification effect. All the non-exploratory results in the preliminary experiment were carefully detailed in a preregistration, and were all replicated in a larger replication experiment, which included the exact same procedure and analysis pipelines. It is important to note that using a larger sample size not only replicated the significance of the effects, but also the descriptive trends, resulting in outcomes of similar effect size.

In an exploratory analysis, we examined variation across stimuli as well as participants, focusing both on the variability in learning patterns between different participants, as well as

within-participant variability in learning for different stimuli. Due to the technical requirements, such an in-depth analysis was applicable only for a smaller subset of our data, however, the partial results we found suggest that the prediction model is applicable in all levels of granularity, both at the individual participant level, as well as in the individual stimulus level.

General discussion

In the current work, we aimed to study and characterize the cognitive mechanisms underlying non-reinforced preference modification with CAT [12], focusing on the training phase itself. The CAT task has a standardized procedure which includes an initial preference evaluation task, a non-reinforced training phase on individual items, and a binary probe phase, the CAT task provided an exceptional opportunity to aggregate together data from multiple different studies and build a large corpus of non-reinforced training data. The current work focused on identifying learning processes via computational modeling of the training data and examining its association with subsequent behavior change.

For this aim, in the first part of the current work we examined previously collected data of n=864 participants from 29 different CAT experiments in a meta-analysis and identified a distinct time-dependent RT pattern during the training task. We found that while in early stages of the training, participants depended on the Go cue to initialize response (homogeneous RTs following cue onset), as training progressed, participants relied less on the Go cue and started to generate fast anticipatory responses. This raised the hypothesis that a marker of transition from cue-dependent responses to early anticipatory responses could be indicative of learning and subsequent preference changes.

Using a Bayesian computational framework implemented with Stan, a statistical model of RT patterns was formulated. RTs were postulated to derive from a mixture model of two Gaussian distributions and a time-dependent mixture proportion, which accounted for the transition speed from late cue-dependent responses to early anticipatory responses. The model parameters were estimated with MCMC algorithm, with a key participant-level θ_{slope_i} parameter, which modeled the individualized difference between participants in transition to anticipatory responses rate. This Bayesian parameter was used as a passive computational marker of learning.

Examining the association between the individualized learning parameter and preference modification effect in the subsequent probe task, showed a positive association in which

711 participants with more robust learning parameter estimates during training also demonstrated
712 stronger preference modification effect in the subsequent probe phase (i.e., enhanced preference
713 for the Go stimuli over NoGo stimuli of similar initial value). Thus the θ_{slope_i} parameter, which
714 was optimize to model RT pattern during training, was found to be good predictor of preference
715 modification effect in a future probe task. An additional analysis modeling RT using a more finely
716 tuned learning $\theta_{slope_{i,s}}$ parameter, fitted for the individual stimuli, further expended these findings,
717 and demonstrated positive predictive power of $\theta_{slope_{i,s}}$ in evaluating preference modification
718 effect on a stimulus-level basis, above and beyond the more general participant-level parameter.

719 Based on the results of Study 1, we derived a theoretical hypothesis with implication for
720 shaping learning efficacy and consequently also the effect of training on preference behavior. If
721 faster transition to anticipatory responses is related to learning, then hindering the ease of cue
722 anticipation could also obstruct learning and resulting in less pronounces preference modification
723 effect. To examine this hypothesis, we design a novel CAT procedure, which was tested in a
724 preliminary experiment and an additional larger preregistered replication experiment. Cue
725 anticipation was manipulated by introducing two training conditions within the CAT task - in the
726 100%-contingency condition, the Go stimuli perfectly anticipated Go cue onset, while in the 50%
727 contingency condition, the Go stimuli only anticipated the Go cue in half of the trials throughout
728 the task.

729 Manipulating cue-contingency during CAT induced the expected effects both in the
730 training and in the probe task. During training, participant relied less on early anticipatory
731 responses and transitioned slower to anticipatory responses in the 50% contingency condition. This
732 effect was captured by the θ_{slope_i} estimates, which were larger in the 100% contingency condition
733 compared with the 50% contingency condition. More importantly, in the subsequent probe phase,
734 a consistent pattern of stronger preference modification effect was observed for the 100%
735 contingency Go stimuli. Thus, manipulating training difficulty has also affected future preference
736 modification measured in the probe phase. When individual differences in preference modification
737 were examined, θ_{slope_i} marker accounted for a large portion of variability in choices, while no
738 significant difference was found between the two contingency conditions above and beyond the
739 θ_{slope_i} effect. This suggest that the more robust preference modification effect observed for the
740 100% contingency Go stimuli, was captured by the stark differences in θ_{slope_i} learning marker.

Furthermore, the fact that intentionally manipulating the training procedure in Study 2 induced differential learning parameter estimates as well as differential behavioral change effect, further supports the hypothesis of a causal nature of the association with the behavioral change. Thus, it is plausible to deduce that shaping training, which was captured by the individualized learning parameter, has driven the induced differential behavior modification effect, as evaluated in the subsequent probe phase. It is important to note that although the cue-contingency was different between the two conditions, the actual exposure-time to the stimuli was identical. Thus, the enhanced preference for Go stimuli in the 100% contingency condition could not be accounted for by mere exposure effect [6].

These findings, showing a consistent link between motor-planning during training and preference modification in probe, correspond with previous works, which showed that rapid response is a crucial feature for preference modification with CAT [13], and neural finding showing that increased striatal and premotor activity during training, were associated with more robust preference modification effect [18]. The results suggest that learning efficacy in the training phase is manifested as stimulus-specific motor planning. Importantly, all participants were alert and attentive during throughout the training phase, as demonstrated by the negligible rates of non-response. However, it evident that mere attention is not sufficient to induce strong preference modification, but rather an attention that can be translated to action, suggesting a unique valuation pathway in the absence of external reinforcements, putatively based on parieto-frontal circuits involving attention with motor-planning [5].

To produce an anticipatory response for a Go stimulus, participants must also rely on memory. Thus, it is possible that the anticipatory response pattern captured in our data identified individualized difference in memory which in turn mediated a change in preferences [22]. This hypothesis is in line with previous findings with CAT which identified a positive association between preferences modification and enhanced memory both at the participant-level [15] and at the stimulus-level [19]. Future imaging studies could examine the new task design with fMRI to identify whether the two contingency conditions can be characterized with differential neural activation pattern.

In addition to a basic understanding of the cognitive mechanisms of non-reinforced learning, identifying a learning marker based on early training runs could be used to design powerful prediction tool for learning efficacy, with relevance to other paradigms and situations

that share the basic training/transfer structure of the current studies. A naïve approach to evaluate learning-efficacy during training tasks such as CAT might be tempted to introduce direct measurements of behavior-change throughout the training process, e.g. by asking participants to make active value-based choices [39,40]. However, it has been shown that the mere act of choice could induce a longitudinal effect of enhanced preference for the chosen stimuli [8,9,41]. Therefore, probing preference within a preference modification training, is likely to alter the learning process and undermine its validity. Moreover, in more standard learning procedures, such as conditioning-based interventions, to examine the efficacy of learning researchers observe the subject's response to the conditioned stimulus when the associated unconditioned stimulus is omitted [e.g., 42–49]. Over repetition of probe tests in all of these procedures might initiate a new learning procedure – where the conditioned stimulus is no longer associated with the unconditioned stimulus, which would eventually lead to extinction-learning [50–53]. Thus, evaluating learning by active probing of the behavior-change effect could interfere with the learning process. However, a passive marker for learning which can predict behavioral change efficacy based on earlier training data, does not evoke the drawbacks of introducing an active choice task to reveal preferences. Using a passive learning marker which is evaluated based on independent training data that preceded the probe phase, overcomes this obstacle. The temporal primacy of θ_{slope_i} estimation could provide a prediction tool for future preference modification in CAT, without tainting the results with a direct evaluating of preferences using a choice task.

The benefits of real-time individualized markers for learning could also be of great value for additional learning and behavior change procedures beyond CAT. In many learning tasks (including CAT), the behavioral change effect on choices is measured in a separate probe phase, proceeding the training phase. For example, in experiments testing pain perception, participants are trained to associate neutral stimuli with differential level of painful heat stimulation before probing the impact of interventions such as drug versus placebo administration [54,55]. Studies examining habit formation may use lengthy free-operant learning protocol, in which participants repeatedly perform an action (such as pressing a button) to gain food rewards, in order to test in a later probe phase whether the participants demonstrate habitual behavior and continue to perform the action even when the reward is devalued (e.g., pressing to get food when satiated) [56–58]. And even outside the field of value-based decision making, in the clinical psychological wellbeing domain, attention bias modification (ABM) procedures use computerized attention training

interventions similar in nature to CAT to treat depression and anxiety [59–61]. In such experimental settings, where training is either unpleasant or exhausting, with plausible odds that learning would not be well-established if under-trained, monitoring learning efficacy individually based on the training data could assist in optimizing training efficacy and efficiency. While the current work focuses on CAT, we assert that the basic mechanisms identified by the current work, could be adapted to accommodate a need in a wider range of learning procedures. With an appropriate adaptation to the desired experimental design, this unique feature of a passive computational marker which is predictive of subsequent change, opens a new path for potential interventions that will allow more efficient training via monitoring and real-time feedback [62,63].

The current work's approach in modeling learning using RT could also be applicable in experimental designs which do not separate the probe from the training task, such as reinforcement-learning tasks, where learning can be evaluated as it progresses on a trial by trial basis [64–67]. Since RT patterns could be indicative of the individual's confidence in her choice during reinforcement learning tasks such as probabilistic selection [48,49] or multi-arm bandit task [68,69], incorporating RT data could improve standard reinforcement learning models and capture learning more accurately [70].

Some limitations were not directly tested in the current work and should be addressed in future studies. Primarily, in Study 2 of the current work, we only demonstrated that learning and preference modification could be hindered using a more difficult association procedure. However, our theory also predicts that easing the association procedure could enhance learning efficacy and preference modification effect. Further work should empirically test this hypothesis, for example by examining the effect of enhancing the Go cue saliency or attention during predefined Go stimuli [25,71,72], or by manipulating the temporal features of training schedule [73]. Furthermore, by instructing participants to respond when they have enough confidence that a Cue will follow, we might have introduced an undesired confound to the task in the subjectivity of confidence each participant need to respond; i.e., some participants might have learned well the association between a stimulus and the Go cue but were reluctant to respond before they validate that in a 50% contingency condition a cue will appear. Although such confound does not undermine the validity of the findings, future studies should avoid it and try to examine a differential effect in which the difference between conditions reflect only task difficulty, for example, by asking to respond to

stimuli which at any point were associated with a cue, or by clarifying in the instruction that a response based on guess is also valid.

Utilizing a Bayesian computational framework, such as the one used here, provided great flexibility required to fit complicated theoretical model. However, the very same advantage of flexibility could also be of disadvantage, when overfitted to the data or abused to examine the same problem from different approaches until reaching a desired outcome, which could have great impact on the reliability and replicability of the results [74–76]. To address this issue, we aimed to use a relatively straightforward model, with as similar features as possible in all studies. We carefully documented the experimental choices in a preregistration before data was collected for an independent replication experiment. Future studies could take the liberty to use the openly accessible data from this work and try to improve the model by changing our current assumptions including distributions' shape, free parameters, and priors.

In conclusion, the current work laid the foundations for understanding and quantifying non-reinforced learning via novel Bayesian modeling approach of RT pattern during training. Using a large meta-analysis dataset of previous CAT studies and two new original experiments, we demonstrated a unique method to evaluate a learning marker with a robust predictive power of subsequent preference change. Thus, we propose that motor response pattern could provide a passive marker for value-change, which does not require direct measurement of preferences.

Methods

Data availability. The entire dataset and analysis codes of the meta-analysis as well as of the novel experiments presented in Study 2 of the current work, are available online at: https://github.com/tomsalomon/CAT_Individualized_Learning.

Study 1 – CAT meta-analysis

Data collection and sample sizes. For study 1 meta-analysis, the data from 29 previous CAT experiment was combined. The dataset included 21 experiments from published works by our research group and collaborators [12–16,18,19,21], as well as eight additional unpublished works, which were collected as part of the preparations for previously published manuscripts, or are planned to be published in the future. The experiments comprised of a median sample size of n = 26 ($M_{sample\ size}$ = 29.79, range = [23, 70]; see Table 1 for information of the sample size in each experiment, and the supplementary data for more detailed description of the unpublished data).

Table 1. **Experiment included in the meta-analysis.**

Exp.	Stimuli	n	Training runs	Go cue	Publication (exp. number in publication [a])
1	Fractal art	25	12	Auditory	Salomon et al., 2018 (2)
2	IAPS positive	27	12	Auditory	Salomon et al., 2018 (3)
3	IAPS negative	28	12	Auditory	Salomon et al., 2018 (4)
4	Snacks (IL)	25	20	Visual	Salomon et al., 2018 (5)
5	Snacks (IL)	25	20	Neg. auditory	Salomon et al., 2018 (6)
6	Faces	25	20	Auditory	Salomon et al., 2018 (7)
7	Fractals	25	20	Auditory	Salomon et al., 2018 (8)
8	IAPS positive	29	20	Visual	Salomon et al., 2018 (9)
9	IAPS negative	32	20	Visual	Salomon et al., 2018 (10)
10	Faces: politic.	25	20	Auditory	Unpublished
11	Faces: politic.	39	20	Auditory	Unpublished
12	Faces	42	16	Auditory	Salomon et al., 2019
13	Faces: affective	42	20	Auditory	Unpublished
14	Faces: affective	70	20	Auditory	Unpublished
15	Fractals	29	16	Auditory	Aridan et al., 2019
16	Snacks (US)	26	12	Visual	Unpublished
17	Snacks (IL)	24	20	Visual (Reward)[b]	Unpublished
18	Snacks (IL)	30	16	Auditory	Botvinik-Nezer, Bakkour, et al., 2021 (1)
19	Snacks (IL)	25	16	Auditory	Botvinik-Nezer, Bakkour, et al., 2021 (Pilot)
20	Snacks (IL)	23	12	Auditory	Unpublished
21	Snacks (IL)	25	20	Auditory	Unpublished
22	Snacks (US)	29	12	Auditory	Schonberg et al., 2014 (1)
23	Snacks (US)	25	8	Auditory	Schonberg et al., 2014 (2)
24	Snacks (US)	25	12	Auditory	Schonberg et al., 2014 (3)
25	Snacks (US)	27	16	Auditory	Schonberg et al., 2014 (4)
26	Snacks (US)	26	16	Auditory	Schonberg et al., 2014 (7)
27	Snacks (US)	30	12	Auditory	Bakkour et al., 2017
28	Snacks (US)	25	16	Auditory	Bakkour et al., 2016
29	Snacks (IL)	36	16	Auditory	Botvinik-Nezer, Salomon, et al., 2019

[a] In papers with multiple experiments, we note in parenthesis the experiment number or title as can be found in the related manuscript.
[b] in experiment 16 a visual cue was used indicating to participants they were awarded with a sum of money, which accumulated throughout the training phase.

All participants gave their informed consent to take part in the experiments. In most experiments, participants received monetary compensation for their time, and in few experiments, some participants took part in the experiment in exchange for course credit. All experiments were approved by the ethical review board of the institutes where they were performed (Tel Aviv University, The Hebrew University of Jerusalem, University of Texas at Austin, and McGill University).

Stimuli. The different experiments included in Study 1 examined the effect of CAT on preferences for various stimuli (see table 1 for a summary of all experiments). In most of the experiments, the stimuli set comprised of images of familiar local snack food items, popular in the US (eight experiments) or in Israel (eight experiments), which participants received for actual

consumption as part of the experiment. Other experiments used face stimuli of unfamiliar figures posing neutral expression from the Siblings dataset [37] (two experiments), unfamiliar faces with neutral and happy expression from the Karolinska directed emotional faces dataset [KDEF; 77] (two experiments) or familiar faces of famous Israeli politicians (two experiments). Unfamiliar abstract stimuli of fractal art [78] were used in three experiments. Two experiments included positive affective stimuli from the international affective picture system [79,IAPS; 80] dataset and two experiments included negative affective stimuli from the IAPS dataset.

The face and snack stimuli were modified using Photoshop to remove any background features and create visual standardization within each experiment. In each experiment, all stimuli were colored images of identical dimensions, where the subject of the image (the snack or the face) was centered in the middle of the image frame and the background was replaced with homogenous background (either black or gray). In experiments with fractal art and IAPS stimuli, the images were only cropped to identical pixel dimensions.

Go stimuli used in the CAT task. In most experiments (23 out of 29) a neutral auditory cue was used as Go cue during the cue-approach training task. In the remaining experiments a neutral visual cue (Experiments 4, 8, 9, and 16), an aversive auditory cue (Experiment 5), or a visual cue indicating a reward (Experiment 17), was used. Both types of visual cues (neutral and reward related) comprised of a semi-transparent Gabor shape, presented on top of the associated Go stimuli. In experiment 17, the appearance of the visual cue indicated to the participant that she or he had won an additional amount of money, which accumulated across the training task.

Procedure. While the different experiments diverged in several aspects of their procedure, all experiments maintained three key phases: an initial-preferences evaluation task, followed by a training task, and a probe task (see Figure 1 for illustration of the procedural design).

Initial preferences evaluation task. In the first task of the experimental procedure, participants were exposed to the complete set of stimuli for the first time and were required to indicate their subjective preferences using one of two tasks. To evaluate participants' preferences in experiments using consumable snack food stimuli, participants performed a Becker-DeGroot-Marschak (BDM) auction procedure [36]. Participants were allocated with either 3 USD (in experiments conducted in the US) or 10 ILS (in experiments conducted in Israel; approximately equivalent to 3.1 USD) which were used to bid on the snack-food stimuli, presented one by one. Participants were informed that at the end of the experiment, one of the trials will be randomly

selected, for which the computer will generate a counter bid. If the participants' bid was higher than that of the computer, they were required to purchase the snack for the lower price bided by the computer. Participants purchased the snack for actual consumption at the end of the experiment. However, if the computer's bid was higher, participants got to keep the allocated sum of money. Participant were explicitly instructed that the best strategy for the task is to indicate their true subjective preference. Prior to their participation, participants were asked to fast for at least 3 hours, to make sure they were hungry and incentivized to purchase the food items, according to their subjective preferences.

In experiments with of non-consumable stimuli (such as fractals and faces), which are less appropriate to be evaluated using a monetary scale, preferences were evaluated using a binary ranking procedure. Participants were presented in each trial a random pair of stimuli and were required to choose the stimuli they prefer better. Based on the idea of choice transitivity, binary choices were quantified to produce subjective value ranks using the Colley Matrix ranking procedure [81].

Following the BDM or binary choice task, stimuli were ranked-ordered according to subjective preferences. The ranks of the stimuli were used as a basis to form two value-groups - one of high-value stimuli (above-median rank) and a group of low-value stimuli (below median rank). The size of the value groups differed between experiments (having 8, 12 or 16 stimuli per value group). In each of the value-groups, half of the stimuli were allocated to be associate with the Go stimuli and response in the subsequent CAT task (Go stimuli), and the other half of stimuli was allocated to be presented in CAT without Go cue (NoGo stimuli).

Allocation for Go and NoGo stimuli within each value category maintained an equal mean rank for Go and NoGo stimuli - e.g., for a high-value group consisting of eight stimuli, the stimuli ranked 8, 11, 12, and 15 were allocated to be Go stimuli, while stimuli ranked 9, 10, 13, 14 were allocated to be NoGo stimuli, such that both allocations were characterized with a mean rank = 11.5. The Go / NoGo allocation was counterbalanced across participants.

Cue Approach Training (CAT). In the CAT task, stimuli were presented individually on the screen center for a fixed duration of one second (except for Experiment 27, where the duration was extended to 1.2 seconds for compatibility with fMRI scanning protocol). In each experiment, all training stimuli were presented once in each training run, thus the number of training runs indicate the number of stimulus repetitions. A fixed proportion of stimuli per experiment

(commonly 30% of stimuli; range 25%-40%) were Go stimuli. When a Go stimulus appeared, a delayed Go cue appeared after a Go signal delay. Participants were asked to respond to the Go cue with a button press as rapidly as possible, before the stimulus offset (one second after the stimulus onset). The Go signal delay was adjusted according to the participant's performance - a failure to respond before stimulus offset resulted in 50ms shortening of the Go signal delay (thus reducing the task difficulty for the following trial), while a successful response on time resulted in 16.667ms increase of the next Go signal delay (making the task more difficult; 1:3 ratio of signal delay increase to decrease). The Go signal delay commonly started at 750ms, and ranged around 700ms (across experiments, $M = 691.78$, $SD = 102.77$).

The training phase in each experiment, consisted of 8 to 20 training runs (see Table 1). Participants were not informed in advance of the contingency between Go signal and Go stimuli. However, as they were repeatedly exposed to the same stimuli, in the later runs of the task, some of the participants were able to identify the Go stimulus and cue association, thus producing accurate faster responses, sometimes even preceding the Go cue onset. This measurement of RT to Go stimuli trials was used in this current work as our main behavioral measurement for the CAT learning model. The details of the model are further discussed below.

Probe. In the final probe phase, preference modification following CAT was evaluated. The probe phase was usually preceded by a 'filler' task such as filling up questionnaires or ranking liking for fractal art images. This task generally provided some time for consolidation and a dissociation between the CAT and probe phases. In the probe task, participants were presented with pairs of stimuli of the similar initial subjective value (both high-value stimuli or both low-value). In each pair, one of the stimuli was a Go stimulus and the other was a NoGo stimulus. Preference modification was evaluated as the proportion of trials in which participants chose the Go stimulus over the NoGo stimulus, above and beyond the expected 50% chance level. This measurement was used as the main outcome variable which indicated preference modification following CAT.

Analysis. In Study 1, we aimed to identify a marker for learning in the CAT task. To achieve this goal, we examined RT patterns in the task. In an exploratory analysis of the meta-analysis data, we identified that as training progressed, mean RT in the task was reduce. We hypothesized that as the training task progresses, participants that were able to identify and learn the stimulus-cue contingency pattern, and thus could generate faster Go responses which do not

rely on the delayed Go cue onset, but rather on the Go stimulus onset. We formally modeled this process in a single experiment before testing it on the entire meta-analysis dataset.

Computational model of CAT. To model our proposed cognitive mechanism we utilized a Bayesian modeling approach of the RT in the CAT task, using the R implementation of Stan programing language [38]. RTs were modeled as a mixture of two gaussian distributions (Equation 1) – one Gaussian of shorter mean RT, representing the early anticipatory responses generated when the stimulus-cue association is predicted by the participants; and a second Gaussian with a later mean RT following the cue-onset (i.e., a distribution representing standard cue-dependent responses).

$$RT_{t,i} - Cue_{t,i} = N(\mu_1, \sigma_{\varepsilon_1})\,(\theta_{t,i}) + N(\mu_2, \sigma_{\varepsilon_2})(1 - \theta_{t,i}) \qquad 1$$
$$\theta_{t,i} = \Phi(-3.1 + \theta_{slope_i} Run_{t,i})$$
$$\theta_{slope_i} \sim N(\theta_{slope}, \sigma_{\theta_{slope}})$$
$$\mu_1 < 0,\ \mu_2 > 0;\ t - trial\ index;\ i - participant\ index$$

The $\theta_{t,i}$ mixture probability of the two Gaussian distribution was used to determine the proportion of trials participants were expected to produce early anticipatory response (1). It was defined using a linear function of time-dependent θ_{slope_i} parameter, which was multiplied by the scaled training run index (training repetition), scaled to 0-1, where 0 and 1 indicating the first and last (20th) training repetition, respectively. This θ_{slope_i} parameter was fitted individually for every participant and was therefore defined conceptually as the individualized learning parameter of interest. To scale the linear function to a range suitable for proportions (0 – 1), we used the normal CDF as a link function.

The baseline proportion of early anticipatory responses (at the first training run) was fixed at the value θ_0 = -3.1, corresponding with probability of 0.1% to generate an anticipatory response. Thus, the individualized learning parameter could be interpreted as the sole parameter effecting the mixture proportion for the two Gaussian distributions, e.g., θ_{slope_i} = 3.1 would be interpreted as a $\Phi(0)$ = 0.5, 50% proportion of anticipatory responses at the final (20th) training run. The individually fitted θ_{slope_i} parameter was determined in the current work as the computational marker of learning and was our main parameter of interest. See Supplementary Code 1 for a complete specification of the model parameters and priors.

The model's parameters were evaluated using Markov-chain Monte Carlo (MCMC) gradient algorithm, implemented with RStan [38]. Each model was evaluated four independent times,

with chains length of 2000 (1000 chain links burn-out time). The mean values of each parameter of the converged model were used as the parameter estimates and are reported along with 95% credible interval (CI). All reported results converged onto stable solution with $\hat{R} = 1$.

To maintain stable interpretation of μ_1 and μ_2 as the centers of anticipatory and cue-dependent responses, respectively, an upper threshold of 150ms from cue onset was imposed on the μ_1 parameter, and a lower threshold of 200ms was for μ_2 parameter. Thus, in every chain μ_1 was imposed the role of the earlier anticipatory responses. Alternative formulation forms of the computational model were also considered and tested, including using 0ms as both upper and lower threshold for the distribution means, using a log-normal distribution instead of a normal distribution, and fitting a unique μ_{1_i} parameter for each participant. However, these models either did not converge when applying to the entire dataset or converged with some issues. Initially we used 0ms as an upper threshold for μ_1, which caused the parameter to converge at this maximal threshold. Using a unique μ_{1_i} parameter for each participant was theoretically favorable (and indeed used in Study 2), but could not be applied in Study 1, potentially either due to computational challenges (fitting many more inter-dependent parameters) or due to the complex structure of the data (e.g., where anticipatory responses were intertwined with late responses due to changing go signal delay)

Computational model of CAT with stimulus-level parameters. In an exploratory model, designed following the analysis of Study 2, we aimed to fit a stimulus-level learning parameter. For each stimulus_s which was presented to participant_i, we fitted an individualized $\theta_{slope_{i,s}}$ parameter, which was used instead of the participants-level individualized learning parameter (θ_{slope_i}), see Supplementary Code 2. Initially, we attempted to model a participant-level dependence, by modeling $\theta_{slope_{i,s}}$ as a parameter derived from higher-level θ_{slope_i} participant-level parameter, similarly to the design in Study 2 (see below). However, such model with tens of thousands of $\theta_{slope_{i,s}}$ parameter estimates that covaried with hundreds of θ_{slope_i} parameters was computationally too demanding. To simplify the model structure, $\theta_{slope_{i,s}}$ parameter was modeled independently of participants' identity (i.e., not considering within-participant possible effect; see Supplementary Code 2).

Deviation from previous version and pre-registration. In our pre-registration, we used an identical Stan model with one key difference – to enforce an order in which μ_1, σ_1 would relate to

early anticipatory RTs and μ_2, σ_2 would relate to late cue-dependent RTs, a different restriction was set in which the upper and lower limit of μ_1 and μ_2, respectively, were set to 0 (Cue onset), instead of the final limits set to 150ms and 200ms. Running the model with these different restrictions converged to a stable solution. Early anticipatory responses were modeled as having an earlier mean (μ_1 = -0.04ms, 95%CI [-0.16, 0.0], σ_{ε_1} = 283.02, 95%CI [280.56, 285.62]) and late cue-dependent responses were very similar to the new model (μ_2 = 287.71ms, 95%CI [287.26, 288.16], σ_{ε_1} = 76.97, 95%CI [76.62, 77.32]). Other parameters were also very similar (θ_{slope} = 3.49, 95%CI [3.20, 3.77]), with variation between participants ($\sigma_{\theta_{slope}}$ = 4.16, 95% CI [3.93, 4.39]). All reported association with probe phase remained very similar.

While the model showed no formal convergence errors (see trace plots of parameters in Supplementary Figure 8), the μ_1 parameter estimate seemed to have been affected by the artificial limitation which was imposed on its maximal value for technical reasons (to avoid inversion with μ_2 across the different chains). Throughout the different chains, the final parameter converged around the upper limit set to 0 (the Cue onset). After several experimental attempts to change this limit, we found the increasing μ_1 lower limit to 150ms resulted in a better solution, in which μ_1 parameter estimate did not converge around the uppermost limit. This model is the final model we chose to use and report here.

Probe analysis. To evaluate the effect of CAT on preferences we analyzed the proportion of trials in which participants chose the Go stimulus over the NoGo stimulus in the probe phase, using mixed-model logistic regression. As Go and NoGo stimuli were matched based on initial value, under the null hypothesis, participants were expected to choose Go stimuli at 50% of trials (log-odds = 0; odds = 1). The results of this analysis were of main interest in previous publications and is reported in the current work for unpublished data (see supplementary materials).

While early work with CAT task showed a differential effect of value on CAT effect on preferences - i.e. CAT usually had induced more prominent preference modification for stimuli of initial high value, compared to stimuli of initial low value [12,13], more recent work with CAT found that this effect was not a dominant feature of CAT [15,18,19,21]. Thus, in the current work, we pooled together data of stimuli with both high- and low-initial value.

Probe and individualized-learning parameter association. Our main analysis of interest aimed to evaluate the association of the CAT preference modification effect with the individualized learning parameter θ_{slope_i}, derived from the computational modeling of CAT. In a

mixed model logistic regression analysis implemented with lme4 R package [82], the number of trial each participant chose the Go versus NoGo stimuli was explained using the independent variable of θ_{slope_i} of each participant, as calculated in the preceding CAT task. The data was aggregated over all participants in the 29 experiments. To account for the aggregation of different experiments in the dataset, we also included random intercept and random slope terms for each of the 29 experiments, see logistic regression formula in Equation 2.

$$Choice \sim 1 + \theta_{slope_i} + \left(1 + \theta_{slope_i} \middle| Exp.\right) \qquad 2$$

In an additional exploratory analysis (which was conceived after the analysis of Study 2), we modeled choices using a stimulus-specific $\theta_{slop\ \ i,s}$ parameter. We aimed to examine if modeling a learning parameter at a more precise stimulus-level (contrary to the pre-registered participant-level $\theta_{slop\ \ i}$ parameter), would serve as a more accurate predictor of future preference modification. For this aim we merged the $\theta_{slope_{i,s}}$ which were estimated for each contingency with the probe data. Choices (number of trials each Go stimulus$_s$ was chosen or not by each participant$_i$) were modeled with two additional models using $\theta_{slope_{i,s}}$ as an independent variable. In the first model, $\theta_{slop\ \ i,s}$ parameter estimates were used instead of θ_{slope_i} estimates. Since $\theta_{slope_{i,s}}$ varied within participants, which were nested within experiment, a nested random slope effect for $\theta_{slope_{i,s}}$ was also added (Equation 3a). In two additional models, probe choices were explained using a full model with both $\theta_{slo\ \ i,s}$ and $\theta_{slop\ \ i}$ (Equation 3b), and a nested model with the same IVs except for the fixed and random effects of $\theta_{slope_{i,s}}$ (Equation 3c). Using a likelihood ratio test for nested models (comparing model 3b with the nested model of 3c), we examined whether $\theta_{slope_{i,s}}$ provided additional explanatory above and beyond $\theta_{slo\ \ i}$.

3

$$Choice \sim 1 + \theta_{slope_{i,s}} + \left(1 + \theta_{slope_{i,s}} \middle| Exp./Participant\right) \qquad \text{(a)}$$

$$Choice \sim 1 + \theta_{slope_{i,s}} + \theta_{slope_i} + \left(1 + \theta_{slope_{i,s}} \middle| Exp./Part.\right) + \left(\theta_{slope_i} \middle| Exp.\right) \qquad \text{(b)}$$

$$Choice \sim 1 + \theta_{slope_i} + \left(1 + \theta_{slope_i} \middle| Exp.\right) \qquad \text{(c)}$$

Effect size estimation for mixed models with R^2_{GLMM}. For the reported logistic regression mixed model, we included an additional effect size score using a generalized linear mixed effect model (GLMM) R^2 estimate, based on the work by Nakagawa, and Schielzeth (2013), implemented with R's Multi-Model Inference (MuMIn) package [83–86]. Like R^2 in linear mixed

model, R^2_{GLMM} was used to quantify the relative proportion of variance accounted by the three generalized mixed model's variance components: variance explained by the fixed effects (σ_f^2), variance explained by the random effects (σ_α^2), and unexplained residual variance (σ_ε^2). In accordance with the developers' suggestion [83,84], we report here two R^2 scores – the marginal R^2 (equation 4a), which signifies the proportion of total variance accounted by the fixed effects (σ_f^2); and the conditional R^2 (equation 4b), which signifies the relative proportion of variance explained by both the fixed and random effects of the model.

4

$$R^2_{GLMM(m)} = \frac{\sigma_f^2}{\sigma_f^2 + \sigma_\alpha^2 + \sigma_\varepsilon^2} \quad \text{(a)}$$

$$R^2_{GLMM(c)} = \frac{\sigma_f^2 + \sigma_\alpha^2}{\sigma_f^2 + \sigma_\alpha^2 + \sigma_\varepsilon^2} \quad \text{(b)}$$

In mixed models examining the contribution of stimulus-level computational marker ($\theta_{slope_{i,s}}$; Equation 3) R^2_{GLMM} values were similarly evaluated. However, it is important to note that the basic unit of analysis in these model (choices per Go stimulus within participant) differ from the unit of analysis in Equation 2 (choices per participant). Thus the R^2_{GLMM} values are not comparable between the two models The $R^2_{GLMM(m)}$ used the participant-level model is the most similar in structure and interpretation to "traditional" R^2 in ordinary linear model, representing the proportion of variance explained by the (fixed) effects of interest, using the participant as the basic unit of analysis.

Study 2: Novel CAT design

Stimuli. In the two novel experiments, the stimuli set included images of 80 unfamiliar faces from the Siblings dataset [37], 40 male and 40 female characters. As in previous CAT studies with this stimuli set [15,18], the images were processed in Photoshop, so that all stimuli were cropped to identical size (400×500 pixels), with the character's pupils positioned at the same spatial coordinates and a homogeneous gray background. The characters poised similar neutral expression and had minimal salient artificial characteristics such as jewelry, make-up, or distinct facial hair. Previous CAT studies with these stimuli found that a similar preferences enhancement for both high- and low-value faces which were associated with a Go cue during CAT [15,18]. Meaning, CAT

had similar effect on preferences both for stimuli of initial low-value and of initial high-value. Using face stimuli allowed us to pool together a larger sample of stimuli of different initial values, under the reasonable assumption that the effect would show similar pattern across the different initial value categories.

During the CAT task, a visual Go cue was associated with some of the face stimuli. The Go cue was identical to the one used in a previous study [15], it comprised of a semi-transparent Gabor image (38×38 pixels, alpha = 0.7), which appeared at the center of the screen, on top of the face stimuli.

Procedure. Like previous CAT experiments, the two novel preliminary and replication experiments included three phases: a baseline preference evaluation task, a modified cue-approach training task, and a post training probe phase which examined preference modification.

Initial preference evaluation task. To evaluate initial preferences, participants underwent a binary forced-choice task between random pairs of stimuli, as in previous CAT experiments with non-consumable stimuli [15,18]. The task included 400 unique choice trials (meaning, no choice between the same two faces repeated more than once), during which each of the 80 face stimuli was presented exactly 10 times. Each choice trial lasted 3000ms, of which participants were given a 2000ms time window to make their choice. Choice trials were followed by a 500ms confirmation screen, showing a green frame around the chosen stimulus, and a fixation cross, which was presented for the remaining trial duration as inter-stimulus interval (ISI), for at least 500ms. In case no choice was made during the 2000ms time-window, a screen saying "You must respond faster" appeared for 500ms, followed by a 500ms ISI.

Binary choices were transformed into individual preference scores using the Colley ranking algorithm [81]. Stimuli were ranked based on each participant's individual initial preferences from the highest value (1) to lowest value (80). Ranks were used to categorize the stimuli into 10 equal-size value groups (each containing eight stimuli; ranks 1-8, ranks 9-16, etc.). The value categories and internal ranks within each category were used to allocate conditions in the subsequent training and probe task, ensuring initial values were balanced across 100% and 50% contingency conditions, as well as across Go and NoGo stimuli within each value-category, which would be pitted against each other in the subsequent probe phase.

Cue-approach training. The CAT task consisted of 20 training runs, in each run all 80 stimuli were presented in a random order for 1000ms each, with a 500ms ISI. In 30% of trials a

visual semi-transparent Go cue appeared 850ms following the stimulus onset for 100ms at the center screen position, on top of the face stimulus. Unlike previous CAT experiments, the Go cue onset did not change throughout the training task. Thus, actual RT was consistent with effective RT (RT from cue onset).

Three Go association conditions were included in this modified version of the CAT – 16 stimuli were always presented with the Go cue (Go stimuli, 100% contingency condition), 16 stimuli were associated with the Go cue during half of the presentations (Go stimuli, 50% contingency condition), and the rest of the stimuli were never followed with a Go cue (NoGo stimuli). Participants were instructed to respond to Go stimuli by pressing a keyboard button as fast as they can, before the face-stimulus disappear. Participant were told in the instruction of the three Go cue contingencies, and that they may respond when they see the or even when they anticipate the cue will appear shortly (see the task instruction as presented to the participants in Supplementary Figure 4). Unlike previous CAT studies, participants were explicitly told of the Go contingency and were thus encouraged to initiate anticipatory responses preceding the actual cue onset, while maintaining high accuracy by only responding when they have sufficient confidence that a cue will follow.

Stimuli were allocated with a go contingency condition based on initial preferences. Stimuli were categorized into 10 equal sized initial value categories, each containing eight stimuli. The stimuli in the highest and lowest initial value categories were allocated to be all NoGo stimuli. Of the middle eight value categories, four were allocated to be 100% contingency and 50% contingency condition category. Within each of the eight middle categories, half of the stimuli were Go stimuli and the other half were NoGo stimuli. The role allocation between categories and within them was designed so that initial values were balanced across the groups E.g., within the second value categories, stimuli ranked 9, 12, 13, and 16 were allocated to be Go stimuli, while stimuli ranked 10, 11, 14, and 15 were allocated to be NoGo stimuli (thus, each group had the same mean rank of 12.5; see illustration in Figure 9). The conditions allocation by initial value was counter-balanced across participants.

Value category	Ranks	Contingency for Go stimuli
1 (highest)	1 – 8	All NoGo
2	9 – 16	100%

Value rank	Go allocation
9	100% Go
10	NoGo

3	17 – 24	50%
4	25 – 32	50%
5	33 – 40	100%
6	41 – 48	100%
7	49 – 56	50%
8	57 – 64	50%
9	65 – 72	100%
10 (lowest)	73 – 80	All NoGo

11	NoGo
12	100% Go
13	100% Go
14	NoGo
15	NoGo
16	100% Go

Figure 9. Go stimuli allocation illustration. Both allocations to Go contingency condition (100% versus 50%) and Go/NoGo stimuli were balanced based on initial subjective value. Go stimuli of 50% contingency were associated in 10 out of the 20 training runs with the Go cue. Different designs were counter-balanced across participants by switching between 100% and 50% contingency categories positions and within category Go/NoGo positions.

All Go stimuli of the 50% contingency condition were associated with the Go cue in the last two runs as well as in eight additional runs (in total - 10 out of 20 run). This was done for potential future implementation of the task in an MRI scanner, which have not been done at the time of this manuscript write-up.

Probe. In the probe phase, preference modification effect was examined using a binary forced choice task, in which participants chose their preferred stimulus of pairs Go versus NoGo stimuli, of similar initial value. Each Go stimulus was pitted against the four NoGo stimuli within the same initial value category (e.g., Go stimuli ranked 9, 12, 13, and 16 were pitted against NoGo ranked 10, 11, 14, and 15). Participants had 1500ms to make their choice, which was followed by a 500ms confirmation feedback (or a message prompting faster response, in case no choice was made), and an ISI of varying duration (1000ms – 9500ms, M = 3000ms), drawn from a truncated exponential distribution with 100ms precision.

Choices of the 100% contingency condition and 50% contingency condition were merged across the different value categories and analyzed with a mixed-logistic regression model. We hypothesized that participants would choose the Go stimuli over NoGo stimuli, and that this preference modification effect would be more robust for the 100% contingency condition. We also hypothesized that individual θ_{slope_i} parameter from computational modeling of the training task, would predict which participants would demonstrate stronger preference modification effect.

Participants. In the preliminary experiment and replication experiment n = 20 and n = 59 valid participants completed the experiment and were included in the analysis, respectively. The sample size of the preliminary experiments was based on the minimal sample size used in previous

CAT studies. The sample size required for the replication sample was based on a power analysis using the results of the preliminary experiment. A sample size of n = 59 was expected to be sufficient to achieve 95% power to detect a significant correlation effect (α = 0.05) between probe choices and the θ_{slope_i} parameter estimates, in both 100% and 50% contingency conditions. The power analysis and the resulting sample sized was documented in the pre-registration and its supplementary materials (https://osf.io/nwr4v).

Three quality-assurance measurements were used as exclusion criteria, based on previous studies with CAT [15,18,21] – (1) low variability of Colley scores in the initial preference evaluation task (which indicate intransitive choice pattern), (2) proportion of false alarm during training, and (3) proportion of missed Go trials during training. In each experiment, we excluded participants with extremely low transitivity, high false alarm, or high miss rate (defined as 3SD from the group mean). The exclusion criteria were pre-registered for the replication experiment, along with the planned sample size.

In the preliminary experiment, no participant was excluded due to the mentioned above exclusion criteria (Transitivity score: M = 0.205, 3SD cutoff = 0.122, min valid score = 0.137; False alarm rate: M = 3.62%, 3SD cutoff = 15.80%, max valid score = 15.27%; Miss rate: M = 6.33%, 3SD cutoff = 24.13%, max valid score = 21.04%). In the replication experiment, five participants were excluded due to these pre-registered exclusion criteria: Two participants had low transitivity score, one participant had high rate of false-alarm, and two participants had high rate of missed Go trials (Transitivity score: M = 0.213, 3SD cutoff = 0.141, M_{valid} = 0.216, min valid score = 0.143; False alarm rate: M = 4.74%, 3SD cutoff = 39.00%, M_{valid} = 3.49%, max valid score = 37.59%; Miss rate: M = 7.59%, 3SD cutoff = 40.22%, M_{valid} = 5.97%, max valid score = 38.75%).

Analysis. Like the procedural design, the analyses of the two experiments were also identical, and generally resembled that of the meta-analysis study, introducing some analysis improvements which could not be applied in the meta-analysis.

CAT computational model. The general goal and design of the Bayesian computational framework in the two new experiments resembled that of Study 1. Participants' RTs were modeled as a mixture of two Gaussian distributions - one distribution of late, cue-dependent responses, and another distribution of earlier anticipatory responses. A $\theta_{t,i}$ mixture proportion determined the probability of making an anticipatory response by the participant$_i$ at trial$_t$. The $\theta_{t,i}$ probability was modeled as a monotonic function of time, using as an individual rate parameter. Unlike the meta-

analysis model, in Study 2, we introduced two distinct contingency conditions, thus for each participant two $\theta_{slop\ \ i}$ parameters were modeled, namely one for the 100% contingency condition, and one for the 50% contingency condition. Another discrepancy between the meta-analysis model and Study 2 model, was an introduction of individual parameter for anticipatory responses center (μ_{1_i}), which allowed flexibility in modeling individual differences in anticipatory responses onsets (Equation 5). This model improvement was possible thanks to the more homogenous nature of the data. Each experiment (preliminary and replication) was modeled separately. See the complete model and priors in Supplementary Code 3.

$$RT_{t,i} = N\left(\mu_{1_i}, \sigma_{\varepsilon_1}\right)\left(\theta_{t,i}\right) + N\left(\mu_2, \sigma_{\varepsilon_2}\right)(1 - \theta_{t,i}) \qquad 5$$
$$\theta_{t,i} = \Phi\left[-3.1 + I_{cond._{t,i}}\theta_{100\%slope_i}Run_{t,i} + \left(1 - I_{cond._{t,i}}\right)\theta_{50\%slope_i}Run_{t,i}\right]$$
$$\theta_{100\%slope_i} \sim N\left(\theta_{100\%,}\sigma_{\theta_{100\%}}\right);\ \theta_{50\%slope_i} \sim N\left(\theta_{50\%,}\sigma_{\theta_{50\%}}\right);\ \mu_{1_i} \sim N(\mu_1, \sigma_{\mu_1})$$

$$\mu_{1_i} < CueOnset + 100,\ \mu_2 > CueOnset;$$
$$t - trial\ index;\ i - participant\ index$$
$$I_{cond._{t,i}} - condition\ index\ (100\%\ versus\ 50\%)$$

In an additional exploratory analysis (not pre-registered), we aimed to create an even more accurate learning computational marker by modeling a $\theta_{slope_{i,s}}$ for every stimulus$_s$ that participant$_i$ was trained with. Assuming that some variability in training speed could be measured not only between participants, but also withing each participant. Since the actual stimuli which were allocated to be Go stimuli varied between participants (i.e., for each participant the condition and role of a certain image stimulus was different), we assumed no mutual information is shared between the same stimulus index of different participants. Stimuli and participants were treated as random effects (Equation 6).

$$RT_{t,i} = N\left(\mu_{1_i}, \sigma_1\right)\left(\theta_{t,i}\right) + N(\mu_2, \sigma_2)(1 - \theta_{t,i}) \qquad 6$$
$$\theta_{t,i} = \Phi\left(-3.1 + I_{stim._{t,i}}\theta_{slope_{i,s}}Run_{t,i}\right)$$
$$\mu_{1_i} < CueOnset,\ \mu_2 > CueOnset;$$
$$t - trial\ index;\ i - participant\ index; s - stimulus\ index$$
$$I_{stim._t} - stimulus\ index\ (for\ each\ contingency\ condition\ separately)$$

In the process of testing this novel approach, the model did not converge when presented with the full data which included both 100% and 50% contingency conditions. However, when trained twice, using each contingency-condition as an independent dataset, the model did converge. See the complete model and priors in Supplementary Code 4.

Probe. As in previous CAT studies, preference modification was evaluated following CAT. Participants' preferences for Go over NoGo stimuli were categorized into two conditions corresponding with the CAT conditions – of 100% contingency and 50% contingency conditions. Trials of the different value groups were pooled together and analyzed with a mixed model logistic regression.

Based on the results of the meta-analysis, we hypothesized that better learning (manifested as larger $\theta_{slop\ \ i}$ parameter estimate) would induce stronger preference modification effect. Thus, we also hypothesized that a stronger preference modification effect would be observed for the 100% contingency condition, compared with the 50% contingency condition. To examine this hypothesis, we run a one-sided mixed model logistic regression with an additional fixed and participant-based random explanatory variables (EVs) of the contingency condition (Equation 7).

$$Choice \sim 1 + Contingency + (1 + Contingency|Participant) \qquad 7$$

Probe choice prediction based on individual learning parameter. Most importantly, we hypothesized that using the θ_{slope_i} parameter estimate for learning would predict future preference modification effect observed during the subsequent probe phase. To test this hypothesis, we introduced the θ_{slope_i} as an additional independent variable in the mixed model logistic regression. Choices were modeled using contingency condition (both as fixed effect and as a random slope between participants), $\theta_{slop\ \ i}$, and an interaction term. This model is equivalent to modeling choices using an intercept and $\theta_{slop\ \ i}$ slope separately for each contingency condition with a random intercept modeled within participants (Equation 8a).

In an additional (not preregistered) exploratory analysis, we modeled a stimulus-specific $\theta_{slop\ \ i,s}$ parameter. We aimed to examine if modeling a learning parameter at a more detailed stimulus-level (extending the pre-registered participant-level $\theta_{slop\ \ i}$ parameter), would serve as a more accurate predictor of future preference modification. For this aim we merged the $\theta_{slo\ \ i,s}$ which were estimated for each contingency with the probe data. Choices were modeled with two additional models using a $\theta_{slope_{i,s}}$ as an independent variable. In the first model, $\theta_{sl\ \ i,s}$ parameter estimates were used instead of $\theta_{slop\ \ i}$ estimates. Since $\theta_{slope_{i,s}}$ varied within participant, a random slope effect for $\theta_{slope_{i,s}}$ was also added (Equation 8b). In another model, probe choices were explained using a full model with both $\theta_{slop\ \ i,s}$ and θ_{slope_i} (Equation 8c). This model was used to

examine whether $\theta_{slope_{i,s}}$ provided additional explanatory power in a likelihood ratio test for nested models (comparing the model of 8c with the nested model of 8a).

$$Choice \sim 1 + I_{cond.} * \theta_{slope_i} + (1 + I_{cond.}|Participant) \quad (8a)$$

$$Choice \sim 1 + I_{cond.} * \theta_{slope_{i,s}} + \left(1 + I_{cond.} * \theta_{slope_{i,s}}\middle|Participant\right) \quad (8b)$$

$$Choice \sim 1 + I_{cond.} * \theta_{slope_i} + I_{cond.} * \theta_{slope_{i,s}} + \left(1 + I_{cond.} * \theta_{slope_{i,s}}\middle|Part.\right) \quad (8c)$$

$$i - participant\ index; s - stimulus\ index$$

$$I_{cond.} - contingency\ condition\ indicator\ IV$$

When results with $\theta_{slope_{i,s}}$ were analyzed, we found that some participants were fitted a $\theta_{slope_{i,s}}$ estimates of extremely small variability (SD<0.1, meaning their $\theta_{slop\ _{i,s}}$ estimates were effectively fixed; see Supplementary Figure 5). In such cases a mixed model with $\theta_{slope_{i,s}}$ random slope (Equation 8b and 8c), resulted in convergence warnings (this phenomenon was not observed in Study 1). Thus, for these analyses, we excluded all participants which demonstrated such low variability of $\theta_{slop\ _{i,s}}$estimates in either one of the two contingency conditions. This resulted in exclusion of nine participants in the preliminary experiment, and 14 participants in the replication experiment (out of 20 and 59 participants, respectively). For the nested model likelihood ratio analysis (comparing a full model to model without $\theta_{slope_{i,s}}$ independent variables), both models were examined using the same subset of 11 and 45 valid participants.

Effect size estimation for mixed models with R^2_{GLMM}. As in the meta-analysis study, we report for the logistic regression mixed models two R^2 estimates – $R^2_{GLMM(m)}$ and $R^2_{GLMM(c)}$, denoting the marginal and conditional R^2 effect (respectively) of the generalized linear model [83–86].

References

1. Staddon, J. E. R. & Cerutti, D. T. Operant Conditioning. *Annual Review of Psychology* **54**, 115–144 (2003).
2. Rangel, A., Camerer, C. & Montague, P. R. A framework for studying the neurobiology of value-based decision making. *Nat. Rev. Neurosci.* **9**, 545–556 (2008).
3. Sutton, R. S. & Barto, A. G. *Reinforcement learning: An introduction*. (MIT press, 2018).
4. Niv, Y. Reinforcement learning in the brain. *J. Math. Psychol.* **53**, 139–154 (2009).

5. Schonberg, T. & Katz, L. N. A Neural Pathway for Nonreinforced Preference Change. *Trends Cogn. Sci.* **24**, 504–514 (2020).
6. Zajonc, R. B. Attitudinal Effects of Mere Exposure. *J. Pers. Soc. Psychol.* **9**, 1–27 (1968).
7. Voigt, K., Murawski, C. & Bode, S. Endogenous formation of preferences: Choices systematically change willingness-to-pay for goods. *J. Exp. Psychol. Learn. Mem. Cogn.* **43**, 1872–1882 (2017).
8. Sharot, T., Velasquez, C. M. & Dolan, R. J. Do decisions shape preference? Evidence from blind choice. *Psychol. Sci.* **21**, 1231–1235 (2010).
9. Sharot, T., Fleming, S. M., Yu, X., Koster, R. & Dolan, R. J. Is Choice-Induced Preference Change Long Lasting? *Psychol. Sci.* **23**, 1123–1129 (2012).
10. Tom, G., Nelson, C., Srzentic, T. & King, R. Mere exposure and the endowment effect on consumer decision making. *J. Psychol. Interdiscip. Appl.* **141**, 117–125 (2007).
11. Rindfleisch, A. & Inman, J. J. Explaining the familiarity-liking relationship: Mere exposure, information availability, or social desirability? *Mark. Lett.* **9**, 5–19 (1998).
12. Schonberg, T. *et al.* Changing value through cued approach: an automatic mechanism of behavior change. *Nat. Neurosci.* **17**, 625–30 (2014).
13. Bakkour, A. *et al.* Mechanisms of choice behavior shift using cue-approach training. *Front. Psychol.* **7**, Article 421 (2016).
14. Bakkour, A., Lewis-Peacock, J. A., Poldrack, R. A. & Schonberg, T. Neural mechanisms of cue-approach training. *Neuroimage* **151**, 92–104 (2017).
15. Salomon, T. *et al.* The cue-approach task as a general mechanism for long term non-reinforced behavioral change. *Sci. Rep.* **8**, article number: 3614, 1-13 (2018).
16. Aridan, N., Pelletier, G., Fellows, L. K. & Schonberg, T. Is ventromedial prefrontal cortex critical for behavior change without external reinforcement? *Neuropsychologia* **124**, 208–215 (2019).
17. Veling, H. *et al.* Training impulsive choices for healthy and sustainable food. *J. Exp. Psychol. Appl.* **23**, 204–215 (2017).
18. Salomon, T., Botvinik-Nezer, R., Oren, S. & Schonberg, T. Enhanced striatal and prefrontal activity is associated with individual differences in nonreinforced preference change for faces. *Hum. Brain Mapp.* **41**, 1043–1060 (2019).
19. Botvinik-Nezer, R., Bakkour, A., Salomon, T., Shohamy, D. & Schonberg, T. Memory for

individual items is related to nonreinforced preference change. *Learn. Mem.* **28**, 348–360 (2021).

20. Zoltak, M. J., Veling, H., Chen, Z. & Holland, R. W. Attention! can choices for low value food over high value food be trained? *Appetite* **124**, 124–132 (2018).
21. Botvinik-Nezer, R., Salomon, T. & Schonberg, T. Enhanced bottom-up and reduced top-down neural mechanisms drive long-lasting non-reinforced behavioral change. *Cereb. Cortex* **30**, 858–874 (2020).
22. Weber, E. U. & Johnson, E. J. Constructing preferences from memory. in *The Construction of Preference* (2009). doi:10.1017/cbo9780511618031.022
23. Weilbächer, R. A., Krajbich, I., Rieskamp, J. & Gluth, S. The influence of visual attention on memory-based preferential choice. *Cognition* **215**, (2021).
24. Kraemer, P. M., Weilbächer, R. A., Mechera-Ostrovsky, T. & Gluth, S. Cognitive and neural principles of a memory bias on preferential choices. *Curr. Res. Neurobiol.* **3**, 100029 (2022).
25. Krajbich, I., Armel, C. & Rangel, A. Visual fixations and the computation and comparison of value in simple choice. *Nat. Neurosci.* **13**, 1292–1298 (2010).
26. Krajbich, I. & Rangel, A. Multialternative drift-diffusion model predicts the relationship between visual fixations and choice in value-based decisions. *Proc. Natl. Acad. Sci. U. S. A.* **108**, 13852–13857 (2011).
27. Krajbich, I. Accounting for attention in sequential sampling models of decision making. *Current Opinion in Psychology* **29**, 6–11 (2019).
28. Ridderinkhof, K. R., Ullsperger, M., Crone, E. A. & Nieuwenhuis, S. The role of the medial frontal cortex in cognitive control. *Science (80-.).* **306**, 443–447 (2004).
29. Tanji, J. New concepts of the supplementary motor area. *Curr. Opin. Neurobiol.* **6**, 782–787 (1996).
30. O'Doherty, J. *et al.* Dissociable roles of ventral and dorsal striatum in instrumental conditioning. *Science (80-.).* **304**, 452–454 (2004).
31. Pessiglione, M., Seymour, B., Flandin, G., Dolan, R. J. & Frith, C. D. Dopamine-dependent prediction errors underpin reward-seeking behaviour in humans. *Nature* **442**, 1042–1045 (2006).
32. Clithero, J. A. & Rangel, A. Informatic parcellation of the network involved in the

computation of subjective value. *Soc. Cogn. Affect. Neurosci.* **9**, 1289–1302 (2014).

33. Kable, J. W. & Glimcher, P. W. The neurobiology of decision: consensus and controversy. *Neuron* **63**, 733–745 (2009).
34. Collins, A. G. E. & Frank, M. J. Opponent actor learning (OpAL): Modeling interactive effects of striatal dopamine on reinforcement learning and choice incentive. *Psychol. Rev.* **121**, 337–366 (2014).
35. Tricomi, E. M., Delgado, M. R. & Fiez, J. A. Modulation of caudate activity by action contingency. *Neuron* **41**, 281–292 (2004).
36. Becker, G. M., DeGroot, M. H. & Marschak, J. Measuring utility by a single-response sequential method. *Behav. Sci.* **9**, 226–232 (1964).
37. Vieira, T. F., Bottino, A., Laurentini, A. & De Simone, M. Detecting siblings in image pairs. *Vis. Comput.* **30**, 1333–1345 (2014).
38. Stan Development Team. RStan: the R interface to Stan. (2020).
39. Hands, D. W. Foundations of Contemporary Revealed Preference Theory. *Erkenntnis 2012 785* **78**, 1081–1108 (2012).
40. Vlaev, I., Chater, N., Stewart, N. & Brown, G. D. A. Does the brain calculate value? *Trends Cogn. Sci.* **15**, 546–554 (2011).
41. Nakamura, K. & Kawabata, H. I Choose, Therefore I Like: Preference for Faces Induced by Arbitrary Choice. *PLoS One* **8**, 1–8 (2013).
42. Barak, S. *et al.* Disruption of alcohol-related memories by mTORC1 inhibition prevents relapse. *Nat. Neurosci.* **16**, 1111–1117 (2013).
43. Joel, D. The signal attenuation rat model of obsessive–compulsive disorder: a review. *Psychopharmacol. 2006 1864* **186**, 487–503 (2006).
44. Tzschentke, T. M. Measuring reward with the conditioned place preference paradigm: a comprehensive review of drug effects, recent progress and new issues. *Prog. Neurobiol.* **56**, 613–672 (1998).
45. LaBar, K. S., Gatenby, J. C., Gore, J. C., LeDoux, J. E. & Phelps, E. A. Human Amygdala Activation during Conditioned Fear Acquisition and Extinction: a Mixed-Trial fMRI Study. *Neuron* **20**, 937–945 (1998).
46. De Houwer, J. Association learning of likes and dislikes: A review of 25 years of research on human evaluative conditioning. *Psychol. Bull.* **127**, 853 (2001).

1394 47. Kawa, A. B., Bentzley, B. S. & Robinson, T. E. Less is more: prolonged intermittent
1395 access cocaine self-administration produces incentive-sensitization and addiction-like
1396 behavior. *Psychopharmacology (Berl).* **233**, 3587–3602 (2016).
1397 48. Frank, M. J., Seeberger, L. C. & O'Reilly, R. C. By carrot or by stick: Cognitive
1398 reinforcement learning in Parkinsonism. *Science (80-.).* **306**, 1940–1943 (2004).
1399 49. Frank, M. J., Moustafa, A. A., Haughey, H. M., Curran, T. & Hutchison, K. E. Genetic
1400 triple dissociation reveals multiple roles for dopamine in reinforcement learning. *Proc.*
1401 *Natl. Acad. Sci. U. S. A.* **104**, 16311–16316 (2007).
1402 50. Calcagnetti, D. J. & Schechter, M. D. Extinction of cocaine-induced place approach in
1403 rats: A validation of the "biased" conditioning procedure. *Brain Res. Bull.* **30**, 695–700
1404 (1993).
1405 51. Bouton, M. E. Context, time, and memory retrieval in the interference paradigms of
1406 Pavlovian learning. *Psychol. Bull.* **114**, 80–99 (1993).
1407 52. Bouton, M. E. Context and behavioral processes in extinction. *Learn. Mem.* **11**, 485–494
1408 (2004).
1409 53. Myers, K. M. & Davis, M. Behavioral and Neural Analysis of Extinction. *Neuron* **36**,
1410 567–584 (2002).
1411 54. Atlas, L. Y., Bolger, N., Lindquist, M. A. & Wager, T. D. Brain mediators of predictive
1412 cue effects on perceived pain. *J. Neurosci.* **30**, 12964–12977 (2010).
1413 55. Koban, L., Kusko, D. & Wager, T. D. Generalization of learned pain modulation depends
1414 on explicit learning. *Acta Psychol. (Amst).* **184**, 75–84 (2018).
1415 56. Pool, E. R. *et al.* Determining the effects of training duration on the behavioral expression
1416 of habitual control in humans: a multi-laboratory investigation. *Learn. Mem.* **29**, 16–28
1417 (2022).
1418 57. Tricomi, E., Balleine, B. W. & O'Doherty, J. P. A specific role for posterior dorsolateral
1419 striatum in human habit learning. *Eur. J. Neurosci.* **29**, 2225–2232 (2009).
1420 58. Dolan, R. J. & Dayan, P. Goals and habits in the brain. *Neuron* **80**, 312–325 (2013).
1421 59. Browning, M., Holmes, E. A., Charles, M., Cowen, P. J. & Harmer, C. J. Using attentional
1422 bias modification as a cognitive vaccine against depression. *Biol. Psychiatry* **72**, 572–579
1423 (2012).
1424 60. Hakamata, Y. *et al.* Attention bias modification treatment: A meta-analysis toward the

establishment of novel treatment for anxiety. *Biol. Psychiatry* **68**, 982–990 (2010).

61. Bar-Haim, Y., Lamy, D., Pergamin, L., Bakermans-Kranenburg, M. J. & van IJzendoorn, M. H. Threat-related attentional bias in anxious and nonanxious individuals: a meta-analytic study. *Psychol. Bull.* **133**, 1–24 (2007).
62. Tempelaar, D. T., Rienties, B. & Giesbers, B. In search for the most informative data for feedback generation: Learning analytics in a data-rich context. *Comput. Human Behav.* **47**, 157–167 (2015).
63. Pardo, A. A feedback model for data-rich learning experiences. *Assess. Eval. High. Educ.* **43**, 428–438 (2018).
64. Leong, Y. C., Radulescu, A., Daniel, R., DeWoskin, V. & Niv, Y. Dynamic Interaction between Reinforcement Learning and Attention in Multidimensional Environments. *Neuron* **93**, 451–463 (2017).
65. Schultz, W., Dayan, P. & Montague, P. R. A neural substrate of prediction and reward. *Science (80-.).* **275**, 1593–1599 (1997).
66. Ratcliff, R. A Theory of Memory Retrieval Roger. *Psychol. Rev.* **85**, 59–108 (1978).
67. Smith, P. L. & Ratcliff, R. Psychology and neurobiology of simple decisions. *Trends Neurosci.* **27**, 161–168 (2004).
68. Niv, Y. *et al.* Reinforcement Learning in Multidimensional Environments Relies on Attention Mechanisms. *J. Neurosci.* **35**, 8145–8157 (2015).
69. Niv, Y., Edlund, J. A., Dayan, P. & O'Doherty, J. P. Neural Prediction Errors Reveal a Risk-Sensitive Reinforcement-Learning Process in the Human Brain. *J. Neurosci.* **32**, 551–562 (2012).
70. Ballard, I. C. & McClure, S. M. Joint modeling of reaction times and choice improves parameter identifiability in reinforcement learning models. *J. Neurosci. Methods* **317**, 37–44 (2019).
71. Shimojo, S., Simion, C., Shimojo, E. & Scheier, C. Gaze bias both reflects and influences preference. *Nat. Neurosci.* **6**, 1317–1322 (2003).
72. Armel, K. C., Beaumel, A. & Rangel, A. Biasing simple choices by manipulating relative visual attention. *Judgm. Decis. Mak.* **3**, 396–403 (2008).
73. Cepeda, N. J., Vul, E., Rohrer, D., Wixted, J. T. & Pashler, H. Spacing effects in learning: A temporal ridgeline of optimal retention: Research article. *Psychol. Sci.* **19**, 1095–1102

(2008).

74. Open Science Collaboration. Estimating the reproducibility of psychological science. *Science (80-.).* **349**, aac4716 (2015).

75. Gelman, A. & Loken, E. *The garden of forking paths: Why multiple comparisons can be a problem, even when there is no "fishing expedition" or "p-hacking" and the research hypothesis was posited ahead of time*. *Department of Statistics, Columbia University* (2013).

76. Camerer, C. F. *et al.* Evaluating the replicability of social science experiments in Nature and Science between 2010 and 2015. *Nature Human Behaviour* **2**, (2018).

77. Lundqvist, D., Flykt, A. & Öhman, A. Karolinska directed emotional faces. *Cogn. Emot.* (1998).

78. Fantastic Fractals. (2013). Available at: http://fantastic-fractals.com.

79. Lang, P. J., Bradley, M. M. & Cuthbert, B. N. *International affective picture system (IAPS): Affective ratings of pictures and instruction manual*. *Technical Report A-8* (2008).

80. Lang, P. J., Bradley, M. M. & Cuthbert, B. N. International Affective Picture System (IAPS): Technical Manual and Affective Ratings. *NIMH Cent. Study Emot. Atten.* 39–58 (1997). doi:10.1027/0269-8803/a000147

81. Colley, W. Colley's bias free college football ranking method: The colley matrix explained. http://www.colleyrankings.com/matrate.pdf (2002).

82. Bates, D., Mächler, M., Bolker, B. M. & Walker, S. C. Fitting linear mixed-effects models using lme4. *J. Stat. Softw.* **67**, 1–48 (2015).

83. Nakagawa, S. & Schielzeth, H. A general and simple method for obtaining R2 from generalized linear mixed-effects models. *Methods Ecol. Evol.* **4**, 133–142 (2013).

84. Nakagawa, S., Johnson, P. C. D. & Schielzeth, H. The coefficient of determination R2 and intra-class correlation coefficient from generalized linear mixed-effects models revisited and expanded. *J. R. Soc. Interface* **14**, (2017).

85. Johnson, P. C. D. Extension of Nakagawa & Schielzeth's R2GLMM to random slopes models. *Methods Ecol. Evol.* **5**, 944–946 (2014).

86. Bartoń, K. MuMIn: Multi-Model Inference. (2020).